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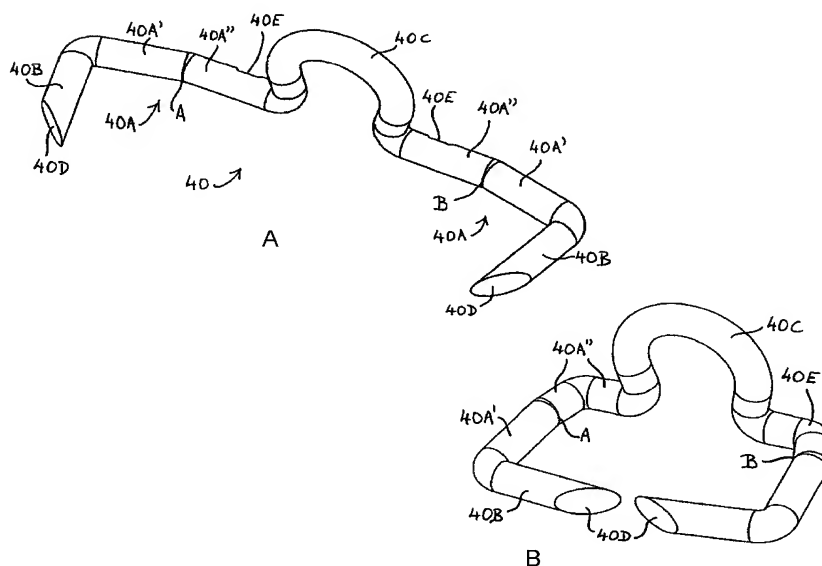
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(54) Title: SURGICAL STAPLE



(57) **Abstract:** A surgical staple (40) comprises a base portion (40A) and a respective leg (40B) extending from each end of the base portion and terminating at a free end (40D), the base portion being deformable to bring the free ends of the legs together to penetrate a liquid-carrying vessel at the site of a puncture and hold the opposite edges of the puncture together. The base portion and legs lie in substantially a common plane except for a centre portion (40C) of the base portion which is deformed in a loop in a direction perpendicular to the common plane. The centre portion of the staple allows the staple to straddle a blood locator tube projecting from the end of a stapler, so that the staple can be closed at the centre of a puncture wound in which the tube is located.



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## SURGICAL STAPLE

### Field of the Invention

5 The present invention relates a surgical staple, for closing a puncture site in a liquid-carrying vessel in a human or animal body, for example a blood vessel, by applying a staple across the puncture so as to affect a closure and prevent bleeding. The invention also  
10 relates to a method of stapling using such a staple.

### Background to the Invention

When performing catheterisation procedures, such as  
15 angiography or angioplasty, a catheter is generally introduced into the vascular system by first penetrating the skin, underlying tissues and blood vessel with a sharpened hollow needle. Next, a guidewire is commonly inserted through the lumen of the  
20 hollow needle and is caused to enter the selected blood vessel. Subsequently the needle is typically stripped off the guidewire and a combination of a dilator and/or introducer (or an introducer alone) are fed over the guidewire and pushed through the skin to enter the  
25 blood vessel. The guidewire can then be removed and a desired catheter to carry out the procedure is fed through the lumen of the introducer and advanced through the vascular system until the working end of the catheter is appropriately positioned. Following  
30 the conclusion of the catheterisation procedure the working catheter will be withdrawn and subsequently the dilator and/or introducer will also be removed from the wound. Following this procedure the vessel puncture

must be closed in order to prevent loss of blood through the puncture hole.

Typically the wound is closed by maintaining external  
5 pressure over the vessel until the puncture naturally seals. This procedure can take approximately 30 minutes with the length of time usually being greater if the patient is hypertensive or anticoagulated. The procedure can also be uncomfortable for the patient and  
10 involves costly professional time on the part of the hospital staff. Other pressure techniques such as pressure bandages, sandbags or clamps have been employed but these also involve ensuring the patient remains motionless for an extended period of time and  
15 is monitored to ensure the effectiveness of the procedure.

A number of devices have been developed in recent times which provide an obstruction in the area of the  
20 puncture in order to prevent bleeding. For example, US Patents 4,852,568 and 4,890,612 disclose a device which utilises a collagen plug which when placed at the blood vessel opening absorbs body fluids, swells and affects a seal. Other plug like devices, for example  
25 US 5,222,974 and US 5,282,827, describe a plug and anchor device, the anchor being positioned inside the vessel and the collagen plug outside the vessel thereby sandwiching the puncture between both and effecting a closure.

30 WO 98/17179 discloses a surgical stapler having a blood locator tube adjacent the stapling head. A guidewire passes through an opening at the end of the tube and up

through a hollow bore in the tube, so that the stapler can be fed onto the guidewire and down onto the puncture site. When the device reaches the puncture site, the tip of the tube enters the blood flow within the artery and blood passes through the tube and out of the distal end at a point visible to the clinician. The clinician can then actuate the stapling mechanism in the knowledge that the stapling head is at the puncture site in the arterial wall.

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It is an object of the present invention to provide an improved surgical staple for closing a puncture in a liquid-carrying vessel. It is a further object of the invention to provide an improved method of stapling using such a staple.

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#### Summary of the Invention

According to the present invention there is provided a surgical staple comprising a base portion and a respective leg extending from each end of the base portion and terminating at a free end, the base portion being deformable to bring the free ends of the legs together to penetrate a liquid-carrying vessel at the site of a puncture and hold the opposite edges of the puncture together, the base portion and legs lying in substantially a common plane except for a centre portion of the base portion which is deformed in a loop at an angle to the common plane.

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In another aspect the invention provides a method of stapling closed a puncture site in a liquid-carrying

vessel in a human or animal body, comprising the steps of:

introducing a stapling mechanism to the location of the vessel;

5 positioning the stapling mechanism at the puncture site by means of an elongated locator device associated with the stapling mechanism, the locator device sensing the position of the puncture site by entering the vessel at the site; and

10 delivering a staple to, and deforming the staple to close, the puncture site;

wherein the staple comprises a base portion and a respective leg extending from each end of the base portion and terminating at a free end, the base portion  
15 being deformable to bring the free ends of the legs together to penetrate a blood vessel at the site of a puncture and hold the opposite edges of the puncture together, the base portion and legs lying in substantially a common plane except for a centre  
20 portion of the base portion which is deformed in a loop at an angle to the common plane; and  
wherein during delivery of the staple to the puncture site the staple loop straddles and slides along the locator device.

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#### **Brief Description of the Drawings**

Embodiments of the invention will now be described, by way of example, with reference to the accompanying  
30 drawings, in which:

Fig. 1 is a perspective view of a surgical stapler incorporating a staple according to the invention;

Fig. 1(A) is an enlarged perspective view of the free end of the shaft of the stapler of Fig. 1;

5 Fig. 2 is a perspective view of the stapler of Fig. 1 with the left-hand side handle removed;

Fig. 3 is a perspective view of the stapler of Fig. 1 with the right hand side handle and shaft removed;

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Fig. 4 is an exploded perspective view of the components seen in Fig. 3 further omitting the left-hand side handle;

15 Fig. 5 is an exploded perspective view of the internal components at the free end of the shaft;

Fig. 6 is a perspective view of the internal components at the free end of the shaft in the pre-fire position  
20 and omitting the left hand side of the shaft;

Fig. 7 is a side elevation of the components of Fig. 6 in the pre-fire position;

25 Fig. 8 is a front elevation of the components of Fig. 6 in the pre-fire position;

Fig. 9 is a perspective view of the internal components of the free end, showing the position of the components  
30 in mid-cycle with fully formed staple;

Fig. 10 is a side elevation of the components of Fig. 9 in the post-fire position;

Fig. 11 is a perspective view of the blood locator tube with enlarged views of the front and rear portions, Fig. 11A and Fig. 11B respectively;

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Fig. 12 is a side sectional elevation of the front portion of an alternative construction of the blood locator tube of the stapler;

10 Fig. 13 is a perspective view of the front portion of the blood locator tube shown in Fig. 12;

Fig 13(A) is a perspective view of the front portion of an alternative construction of the blood locator tube  
15 shown in Fig. 12;

Fig. 14(A) is a perspective view of the surgical staple in the pre-fire (pre-deformed) state;

20 Fig. 14(B) is a perspective view of the surgical staple in the post-fire (deformed) state;

Fig. 15 is an enlarged perspective view of the cam mechanism;

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Fig. 16 is a side elevation of the cam mechanism;

Fig. 17 is a side elevation of the shaft section of the device and suction port;

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Fig. 18 is an end view of the surgical staple, locator tube and insert;

Figs. 19(A) to (C) are perspective views of the staple of Fig. 14 showing alternative locations for bruise points;

- 5 Figs. 20(A) and (B) are pre- and post-fire views of a second embodiment of staple according to the invention;

Figs. 21(A) and (B) are pre- and post-fire views of a third embodiment of staple according to the invention;

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Figs. 22(A) and (B) are pre- and post-fire views of a fourth embodiment of staple according to the invention;

- 15 Figs. 23(A) and (B) are pre- and post-fire views of a fifth embodiment of staple according to the invention; and

Fig. 24) is a pre-view of a sixth embodiment of staple according to the invention.

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#### **Detailed Description of the Preferred Embodiments**

- Referring to the drawings, the stapler comprises a rigid shaft 10 extending from a moulded plastic housing 12 shaped in the form of a pistol-like handle. The shaft 10, which is hollow to accommodate various moving components to be described, comprises right and left hand sides 10A, 10B respectively which are secured together at the distal free end by a section of heat shrinkable tubing 91 in combination with interference pins and mating cavities 15A and 15B (Figs. 4 and 5) along the edges of the distal tip, and at the proximal end by pins 17A mating in an interference fit with
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corresponding cavities 17B (Fig. 2 and 3) captured within the housing 12. Likewise, the housing 12 comprises left and right-hand sides 12A, 12B respectively.

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The major part of the exposed length of the shaft 10 has a constant circular cross-section, but at its free end the shaft 10 has a portion 14 of increased diameter having a "bullet" profile. One end of this bullet  
10 portion 14 is tapered down toward a staple exit slot 16 while the other end is tapered down to the remaining section of the shaft, which extends back into the housing 12. The ratio of the maximum diameter of the bullet portion 14 to the diameter of the remaining  
15 section of exposed shaft is approximately 5:4. Heat shrink sleeve 91 sits flush with the surface of the bullet portion 14, to ensure atraumatic entry, percutaneously, into the tissue.

20 The reason for the bullet profile is so that the shaft 10 is as atraumatic as possible during introduction to the body to minimise the amount of force and tissue dilation required when tracking the device percutaneously over a guidewire 18 and onto the surface  
25 of a blood vessel adjacent a puncture hole, as will be described. In an alternative construction, not shown, the bullet portion 14 is oval in cross-section with the major axis of the oval being coincident with the staple exit slot 16, so as to minimise the circumferential  
30 length for a given staple width.

The bullet portion 14 of the shaft 10 houses a staple 40 and a staple delivery mechanism (Figs. 4 to 7). The

staple delivery mechanism comprises a tiltable anvil 24 and a pair of rod-like actuating members, namely an elongated anvil support 30 and an elongated staple former 52, the latter being slidable in the shaft 10 and operated by a trigger-operated cam mechanism 62 in the handle housing 12.

The anvil 24 has a pair of upstanding fingers 24A at the front and a pair of downwardly inclined tilt arms 24B at the rear. The anvil 24 is tiltably mounted in the bullet portion 14 by a pair of wings 26 which are pivotable in recesses 28 in the right hand side 10A of the shaft 10 (the wings 26 are retained in the recesses by the underside of projections 54 on the former 52).

Tilting of the anvil 24 is effected by the cam mechanism 62 via the anvil support 30, which is slidable axially within the right hand shaft side 10A in channel 32. The front end of the anvil support 30 is bifurcated to form two arms 34 having lateral projections 36 (Figs. 6 and 7). The arms slide in rebates 38 in the right hand shaft side 10A. The anvil support 30 is movable, by the cam mechanism 62, from a forward position, Figs. 6 and 7, wherein the arms 34 extend under the anvil's support wings 25 to support the anvil forming fingers 24A directly in front of a surgical staple 40 to be delivered, to a rearward position, Fig. 10, wherein the arms 34 are withdrawn under the downwardly inclined tilt arms 24B at the rear of the anvil 24 so as to tilt the anvil anti-clockwise (as seen in Fig. 10) and displace the fingers 24A out of the path of the staple 40.

Referring additionally to Figs. 11, 11A and 11B, a hollow blood locator tube 92 is slidable axially within the shaft 10 in a channel 44 in the anvil support 30 and in an opposing U-shaped channel 53 in the staple former 52. The tube 92 extends the full length of the shaft 10 and has a constant, generally oval or elongated cross-section, except at its distal tip 14 where the locator tube 92 is formed into a narrow opening 96 and at a crimped region 94 towards the rear of the tube 92 which is formed to allow only the guidewire 18 and not blood to exit the rear of the locator tube.

Under the action of the cam mechanism 62 the tube 92 is slidable axially in the shaft 10 between a forward position, Figs. 6 and 7, wherein its front end projects beyond the bullet portion 14 of the shaft 10 under the influence of a leaf spring 88 to be described, and a rearward position, Figs. 9 and 10, wherein the front end of the tube 92 is retracted within the bullet portion 14 behind the fingers 24A of the anvil 24 during the rotation of cam 62.

The purpose of the blood locator tube 92 is to follow a previously placed guidewire 18 to a puncture site in a blood vessel, thereby to locate the free end of bullet portion 14 of the shaft 10 against the exterior wall of the blood vessel at the puncture site. To properly locate the bullet portion 14 the front end of the tube 92 must actually penetrate the blood vessel wall at the puncture site and this is indicated by blood flowing back through the tube 92 and out through a blood outlet port 93 (Fig. 11) in the tube. A channel (not shown)

in the part of the left hand side 10B of the shaft 10 within the housing 12 provides communication between the port 93 and a blood exit port 50 (Fig. 1) on the side of the housing 12B, so that the blood flowing back  
5 through the tube 92 is visible at the exterior of the housing.

A blood exit port adapter 51 (Fig. 1) may be secured into the opening of the blood exit port 50 via a  
10 matching male luer taper 51A to enhance the visibility of the exiting blood. The blood exit port adapter has a reduced internal diameter, relative to the opening of the blood exit port 50, which for a constant blood flow increases the pressure of exiting blood causing a jet  
15 effect of exiting blood.

In the absence of the blood exit port adapter, the blood exit port's female luer taper opening matches that of the standard medical syringe's male luer taper  
20 making it possible at any time during the device's use to inject fluid via the blood exit port into the lumen of the locator tube to exit at its distal tip. This may be necessary from time to time to clear the locator tube's lumen of congealed blood or trapped soft tissue.  
25 Alternatively, radiopaque contrast medium may be injected via the locator tube to confirm the relative location of the locator tube's distal tip to that of the blood vessel wall by fluoroscopy, or any injectable fluids may be injected for diagnostic or therapeutic  
30 reasons.

The blood outlet port 93 is sized to have a minimum area corresponding to the available blood entry area at

the distal tip; however, is narrower (in a transverse aspect) than the diameter of the guidewire 18 to prevent the guidewire inadvertently exiting the blood outlet port during insertion, instead of exiting from  
5 the intended proximal end of the locator tube.

It has been found that the naturally formed shape of puncture wounds in arterial walls is elongated rather than round. Whereas the hole is formed by introducing  
10 instruments generally of round cross section, the wall tends to open generally along a transverse line which lies in the direction of the circumference of the artery (rather than along the axis of the artery). By having a generally oval blood locator tube, the locator  
15 tube (when introduced by the clinician with the major axis of the oval perpendicular to the axis of the artery), will fit more naturally within the arterial opening. The consequence of this is that the wound edges which are to be stapled together, lie closer  
20 together than if a tube of circular cross section were to be used.

This in turn has the consequence that the staple used need not be so large, and in turn, the dimensions of  
25 the shaft, which must accommodate the staple when in its unformed state, can be reduced, leading to less trauma for the tissue into and from which the shaft is introduced.

30 A further consequence of having a generally oval or elongated cross section for the locator tube is that the tube will be more disposed to the centre of the puncture than with a rounded tube. The present

embodiment has a staple which straddles the locator tube, thereby increasing the likelihood of the staple closing the elongated wound at its centre rather than towards one or other of the extremities of the wound.

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The opening 96 at the front of the tube 92 has an approximately circular portion 96A at the extreme forward tip of the tube which is of greater diameter than the width of the remaining portion 96B of the opening 96. The portion 96B is in the form of a slot which is aligned with the major axis of the elongated cross-section of the tube 92 and slopes rearwardly from the circular portion 96A. The guidewire 18, which passes through the tube 92, Fig. 11, is chosen to be of sufficiently smaller diameter than the diameter of the opening 96A at the front end of the tube 92 for the guidewire 18 to be easily inserted into the tube 92 and pass through the opening 96A. However, the guidewire is also chosen to be too large to fit within the remainder 96B of the opening 96. In this way guidewire 18 is constrained to remain in opening 96A, and the size of opening 96A sets an upper limit on the diameter of guidewire which can be used with the device. One could introduce a narrow neck or constriction into the opening 96 just above opening 96A (at the points indicated by 96C) to ensure that very small guidewires were constrained within the enlarged opening 96A, but in general this is unnecessary as the guidewire will normally be supplied with the device, or the device will only be supplied for use with a particular gauge of guidewire.

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- The rear crimp 94 and tip opening 96A are positioned to encourage the guidewire to lie along the bottom curved surface of the tube, i.e. that portion of the tube lying in a direct line between the opening in the crimped end and the opening 96A. This helps prevent guidewire 18 from laying up against the inside of blood exit port 93 and preventing egress of blood, Fig. 11A and 11B.
- 10 The curvilinear nature of opening 96 increases the available inlet area to match that of the available area within the body of the locator tube with the guidewire 18 in situ.
- 15 The slot-like opening 96B slopes away from the circular opening 96A for ease of insertion into the vessel opening and to reduce the potential of trauma to the inner wall of the vessel opposite the opening being stapled. This is achieved because the guidewire 18 protruding from opening 96A will tend to push the opposite wall of the vessel away from the locator tube tip, and the point at which the guidewire protrudes (due to it being constrained in the opening 96A) is the farthest part forward of the tip. Thus, the shape of the tip is streamlined away from opening 96A to prevent any part of the tip gouging into or otherwise damaging the inner vessel walls. Also, the peripheral edges 95 of the opening 96 are bent inwardly to as to avoid sharp edges which might damage soft tissue and the vessel wall.
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The distal end of an alternative construction of a locator tube 42 is shown in Figs. 12 and 13. This

construction also has a substantially constant elongated cross-section, which in this case converges to an approximately circular guidewire opening 46 at the extreme forward tip of the tube. The guidewire 18, 5 which passes through the tube 42, is usually chosen to be of sufficiently smaller diameter than the diameter of the opening 46 for there to be an adequate gap for the blood to pass back through the tube 42 even in the presence of the guidewire. However, further openings 10 46A are provided in opposite sides of the tube 42 just behind the front opening 46 to allow more ready access of the blood to the interior of the tube in cases where the guidewire 18 may not leave a large enough gap for passage of blood solely through the opening 46. The 15 three openings 46, 46A, 46A in fact form respective portions of a single front opening, being in reality three connected lobes, all connected by constricted channels 47, and all in communication with the interior of the tube.

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An alternative construction is shown in Fig. 13(A) where the three openings 46, 46A and 46A, while collectively constituting the front opening of the tube 42, are independent of each other. Again, opening 46 25 at the front of the tube is sized to receive a maximum size of guidewire and openings 46A are sized to allow a sufficient flow of blood to enter the locator tube.

A problem can arise in devices of this type where an 30 oversized guidewire is used which occludes the hollow interior of the blood locator tube and thereby prevents blood flow back through the tube. To prevent this situation the lobe 46 through which the guidewire

emerges in the tip of the tube of Figs. 12, 13 and 13A is of a lesser diameter than the internal bore of the tube. The dimensions of this lobe 46 set a maximum for the guidewire diameter for use with the device, and  
5 ensure that even when this maximum diameter guidewire is used, there is still sufficient internal clearance within the tube bore to allow a strong blood flow through the tube from the other lobes 46A.

10 The staple 40 straddles the blood locator tube 92 within the bullet portion 14 of the shaft 10, see Figs. 6 and 8, and is slidable thereon forwardly towards the free end of the bullet portion 14. In particular (see also the enlarged view of Fig. 14), the staple 40  
15 comprises a back or base portion 40A from which extend approximately perpendicularly at each end respective legs 40B which terminate in sharpened points 40D. The legs 40B preferably converge at a small angle of, for example, about  $10^{\circ}$ - $15^{\circ}$ .

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The base portion 40A and legs 40B lie in substantially a common plane except for a centre portion of the base portion 40A which is deformed in a direction perpendicular to the legs 40B so as to have an  $\Omega$

25 (omega) shaped loop 40C generally complementary to the external cross-sectional profile of the blood locator tube 92 and internal cross-section of an insert 160 to be described.

30 At points A and B on each side of the omega section 40C, and at an equal distance therefrom, the outer ends 40A' of the base section 40A are pre-bent forwardly to make an angle of between  $150^{\circ}$  and  $170^{\circ}$  with the inner

ends 40A" which lie substantially on a straight line, the pre-bend points A and B being positioned to maximise the closure of the closed staple (and is relevant to the depth of forming wings 54 on the former 5 52). The base section is also locally deformed ("bruised") at points 40E so as to narrow the cross sectional width of the wire at both points thereby directing the staple to bend at these points. As seen in Fig. 19, the bruise points 40E may be provided on 10 the front of the base section (Fig. 19A), on the rear of the base section (Fig. 19B), or on both the front and rear of the base section (Fig. 19C).

The tips of the points 40D are directed inwardly, i.e. 15 in the direction in which the legs 40B are closed in use, as shown in Fig. 14B. Alternatively, the point geometry may involve two bevel planes so as to create a sharper pointed end (Fig. 24). The point geometry, angles of projecting legs 40B to base 40A and the 20 length of legs 40B are configured to aid in keeping the closing staple within the thickness of the arterial wall and prevent the staple-points 40D penetrating into the arterial lumen, and additionally to avoid snagging on or dislodging any atheromatous or calcified plaque 25 between the intimal and endothelial layers of the arterial wall.

The staple 40 is mounted on the blood locator tube 92 such that the centre portion 40C of the staple sits on 30 the upper half of the tube 92, as seen in Fig. 6 and 8, where the narrow open section of the omega shape is approximately equal to the width of the tube and with the legs 40B pointing forwardly on opposite sides of

the tube 92. The depth of the centre portion 40C of the staple 40 is such that the legs 40B of the staple lie substantially directly on opposite sides of the central axis of the tube 92. This will ensure that the staple 40 is positioned centrally across the puncture hole in the blood vessel. In order to avoid the guidewire 18 fouling the staple 40 when the latter is closed on the puncture site, the hole 96A is offset below the plane containing the legs 40B of the staple, Fig. 8.

The metal insert 160 is received in a recess in the left-hand shaft side 10B within the bullet section 14. The insert 160 provides mechanical support for the omega section 40C of the staple 40 during the staple forming process and is engaged by the former 52 during the staple ejection phase of the process so as to separate both halves of the bullet section for easy staple release. The insert is profiled to generally correspond with the external profile of the omega shaped portion 40C of the staple. At the distal end the insert profile tapers down to closely approximate the omega-shaped portion of the staple 40C (Fig. 18) This has the effect of offering mechanical support to the omega-shaped portion of the staple during the staple forming process, during which the base section is bent about the anvil fingers 24A at the bruise points 40E. This bending motion in turn causes the omega to open up or flatten out. The metal insert prevents this from happening, only allowing the staple base to deform around the anvil at the bruise points 40E. The omega interlock system between the staple 40 and insert 160 (Fig. 18) also stabilises the staple,

vertically, within the staple exit plain during the forming process, whilst allowing easy staple release once formed, due to the relatively small contact area between staple and insert.

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The staple former 52 has a cross-section conforming to that of the blood locator tube 92 and is slidable on the blood locator tube 92 axially within the shaft 10. The former 52 is located behind the staple 40 on the tube 92 and is operated by the cam mechanism 62. At its front end the former 52 has a pair of forming arms 54 which are so shaped that, when the former 52 is driven forward by the cam mechanism 62, the staple 40 is driven against and deformed around the anvil fingers 24A so that the legs 40B of the staple close together (Fig. 9) onto the puncture site. The surface of the forming arms which contact the staple 55 may be so profiled to match the cross-sectional geometry of the staple. This matching profile stabilises the staple on the forming surfaces of the forming arms 54 during the high pressure contact with the staple during staple forming and closure. During the forward movement of the staple, the staple legs slide toward the anvil 24 along a track defined by the staple exit slot 16 between the opposite halves the bullet portion 14. The slot 16 provides a slight interference fit on the staple legs 40B to prevent the staple 40 moving forward during storage of the device or prior to firing. The slot 16 further prevents the staple rotating in the horizontal plane (Fig. 7 and 10) during its forward travel. Once forming of the staple around the anvil is completed the forming force is removed from the former 52 by a drop-off in the cam, the anvil is lowered and

the former advanced again to eject the staple from the device. During this forward movement (ejection phase), the sloped edges 52A and 52B of the former engage with the metal insert 160 to prise open the bullet section  
5 of the shaft assembly thus facilitating staple release.

The cam mechanism 62 can be seen in Fig. 3 and in enlarged views of Figs. 15 and 16. The mechanism 62 consists of a first cam 58 and a second cam 60 mounted  
10 on a common axis 62 which sits in a recess 64 in the left-hand side 10A of the shaft (Fig. 4) and a corresponding recess (not shown) in the right-hand side 10B. Trigger 56 is similarly mounted in the shaft by a pair of stub axles 66 which are received in a trigger  
15 seating recess 68 in each half of the shaft 10, Fig. 4.

An actuating pin 70 extends through the first and second cams 58, 60. This actuating pin is acted on by a cam actuating surface 72 (Fig. 3) provided on the  
20 trigger 56, so that when the trigger is squeezed the actuating surface moves the actuating pin in an anticlockwise direction around the axis 62. Because the actuating pin extends through both cams 58, 60 of the mechanism 62, the cams are both rotated  
25 simultaneously through the same angle as determined by the trigger squeeze. The use of this cam mechanism ensures accurate timing and positive mechanical displacements of all the moving components and accurate movement of the components relative to each other. The  
30 geometry of the trigger pivot pins 66 and actuating surface 72 relative to the cam pivot 62 and cam actuating pin 70 is configured to minimise the trigger rotation to only 23 degrees whilst the cam rotates a

total of 90 degrees. This configuration also provides a mechanical advantage that the trigger delivers to the cam-actuating pin 70 of approximately 1:4. This geometry is further configured to deliver the best  
5 mechanical advantage at the phase during the staple forming cycle, which requires the highest forming forces, having the advantage of minimising the trigger effort and ensuring a constant trigger effort over the full cycle. Trigger 56 further comprises a ratchet  
10 lever 73B, shown in Fig. 3, which engages with ratchet strip 73A, which is mounted in the right handle 12A, Fig. 3. This non-return ratchet system ensures the firing cycle of the staple is uninterrupted, non-repeatable and provides a positive indication that the  
15 device has been used.

Referring back to Fig. 3, a leaf spring 88 positioned in a recess in the left-hand side 10A of the shaft and a corresponding recess (not shown) in the right hand  
20 side 10B. The free ends of the spring are formed into a loop so as to pivot freely in the curved corner recesses in which it sits and to aid assembly. The apex of this spring is positioned in a slot 74 in the crimped portion 94 of the blood locator tube 92 thus  
25 assuming the role of cam follower for the blood locator tube. This blood locator tube cam follower 74 is acted on by the first cam 58. Similarly, the first cam 58 acts on a former cam follower 76, whereas the second cam 60 acts on anvil-support cam followers 78A and 78B.  
30 The shape of the first and second cams 58, 60 are shown in elevation in Fig. 16 (the second cam 60 is shown in dotted outline as it is concealed by the first cam). Fig. 16 also shows actuating pin 70, and a reinforcing

strut 80 mounted between the first and second cams diametrically opposite the actuating pin 70.

5 The cams are shown in the starting positions in Fig. 15 and 16. Squeezing the trigger fully (through an angle of 23 degrees) causes the cams to rotate anticlockwise through 90 degrees.

10 The apex of the leaf spring 88 which engages with and operates as a cam follower for the blood locator tube (leaf spring apex) acts against the rear surface 82 of the first cam 58. As the first cam rotates anticlockwise from the position shown in Fig. 15, the distance between the blood locator tube cam follower 74  
15 and the axis 62 is increased. This causes the blood locator tube to be drawn backwards as the trigger is squeezed.

The former cam follower 76 acts against the front  
20 surface 84 of the first cam 58. Again the distance between former cam follower 76 and axis 62 increases through the initial stages of the trigger being squeezed. The profile of surface 84 is designed with two distinct non-linear efficiencies, transitioned from  
25 low mechanical efficiency/high displacement to high mechanical efficiency/low displacement. The first rise rate being for displacement of the staple from its starting position to initial forming against the anvil, which requires the largest displacement of the staple  
30 with minimal load. The second non-linear rise rate is designed to correlate the cams mechanical efficiency with the load profile required to form the closed staple, minimising the trigger effort required and

ensuring a constant trigger effort over the full cycle. A V-shaped section 84A of front section 84 causes the former 52 to momentarily suspend its forward motion when the staple has been fully formed. The effect of  
5 this is to momentarily release the pressure off the formed staple against the anvil, allowing the anvil to be dropped. The geometry of the distal tip of the former is designed to provide sufficient intrinsic spring tension to allow the forming arms 54 to further  
10 squeeze the formed staple, once the anvil has dropped, to further closed the formed staple. As the cam continues to rotate the raised profile 84B on the cam causes the former to advance forward again, ejecting the staple clear of the device.

15 It can be seen that a raised hump 82A on the profile of the rear surface 82 of the first cam is located almost diametrically opposite the V-shaped section 84A. The reason for this is to increase the rate at which the  
20 blood locator tube is drawn out of the puncture site just before the staple is fully formed and released. The intention is to leave the tube in the puncture as late as possible to provide support for the walls of the blood vessel for as long as possible And also to  
25 ensure that the head of the device remains centred over the puncture hole. The blood locator tube 92 is biased forward by the blood locator tube leaf spring 88 which also maintains pressure between the apex of the spring and the rear surface 82 of the first cam 58.

30 The blood locator tube leaf spring 88 allows the locator tube to be displaced in a proximal direction (back into the shaft of the device) against the spring

tension in the event that the locator tube meets any significant resistance during insertion of the device, to prevent unnecessary trauma to soft tissues, the vessel or its rear wall.

5

An example of where this is particularly useful is if the stapler is advanced too far into the vessel, so that the tip of the tube 92 meets the inner wall. The blood locator tube will then be displaced back into the shaft, and may be designed to protrude through the end of the handle housing to give a visual indication that

10

the device has been inserted against the wall. Furthermore, the device may be designed so that the blood outlet port 93 on the tube 92 is brought out of registry with the blood exit port 50 in the handle housing when the tube is displaced backwards, so that the clinician will note the flow of blood ceasing when the tube meets the inner vessel wall in this way.

15

20 The cam mechanism 62, however, provides positive mechanical displacements for withdrawing the locator tube at the appropriate timing, to ensure there is no chance of the staple being formed whilst the locator tube is in a forward position and potentially

25

A further reason to leave the blood locator tube in the puncture hole as late as possible is that the continued retraction of the tube everts or turns outwards the opposed edges of the puncture wound and aids penetration of the staple legs into the arterial wall. Eversion of the edges of the puncture helps prevent thrombus formation within the vessel. Yet another

30

- reason to leave the blood locator tube in the puncture hole as late as possible is to ensure that the stapler head remains centred over the hole during the staple delivery process. When the locator tube is fully
- 5 retracted, only the guidewire is left within the wound, and this will be easily retracted from the closed wound after the stapler has been removed from the puncture site.
- 10 The anvil-support cam follower 78B acts against the rear surface 90 of the second cam 60. It can be seen that this rear surface 90 provides the greatest increase in distance relative to the axis to the
- 15 horizontal. The reason for this is that the anvil is maintained in place until the staple has been formed and the pressure on the former has been relaxed slightly to allow the anvil to drop. The anvil is maintained in place for the initial 60 degrees of
- 20 rotation by the anvil-support cam follower 78A being in contact with cam surface 98 of cam 60, preventing the anvil-support 30 from moving from its starting position. The cam surface 98 for the first 60 degrees of cam rotation is at a constant distance from the cam
- 25 axle 62 (in dwell).

In use, the stapler is initially in the "pre-fire" configuration shown in Figs. 6 to 8. The front end of the blood locator tube 92 is in a fully forward

30 position projecting beyond the free end of the bullet portion 14 of the shaft 10, the anvil-support 30 is in a fully forward position with its arms 36 extending under the anvil's support wings 25 ensuring the anvil

fingers 24A are directly in front of the staple 40, the former 52 is in a fully retracted position away from the anvil fingers 24A, and the staple 40 is in its fully back position up against the forming arms 54.

5

In this configuration the external end of a previously positioned guidewire 18 is inserted into the hole 96A in the front end of the blood locator tube 92 and fed through the tube 92 until it exits a guidewire exit port at the rear of the housing 12. The stapler is now fed along the guidewire 18 until the tip 95 of the tube 92 enters the blood vessel lumen through the vessel's puncture hole. This is indicated by blood flowing out of the blood exit port 50 or, if present, the adapter 51. At this point the front end of the bullet portion 14 of the shaft 10 will be resting against the exterior wall of the blood vessel.

Now the trigger 56 is squeezed, causing the cams of the cam mechanism 62 to rotate through 90 degrees. As mentioned, the rear end of each of the blood locator tube 92, anvil-support 30 and former 52 are coupled to the cam mechanism via cam followers and the following co-ordinated movement of these components takes place as the cams rotate through 90 degrees.

(A) .

**0 degrees:** Stapler in pre-fire configuration.

**32 degrees:** Former 52 forward sufficiently to clamp staple against anvil fingers 24A, blood locator tube begins to retract. At this point the staple legs will have punctured the wall of blood vessel, but the staple is not yet fully deformed.

**50 degrees:** Former 52 forward sufficiently to deform the staple legs around the anvil fingers 24A and close the staple on the puncture site: blood locator tube 42 fully retracted. At some point between 32 and 50

5 degrees, the blood locator tube will have withdrawn from between the staple legs in time to allow them to close. This should be left as late as possible to provide support for the walls of the blood vessel for as long as possible.

10 **65 degrees:** Clamp force released from staple (due to drop off in cam profile). Anvil support 30 starting to retract.

**75 degrees:** Anvil support 30 retracted sufficiently to act against anvil sloped tilt arms 24B. Anvil fingers

15 24A begin to drop.

**83 degrees:** Anvil support 30 fully retracted. Anvil fingers 24A dropped down to allow release of staple. Intrinsic tension in former arms 54 further closes the staple. Former 52 begins to move forward again to

20 eject staple. Former 52 begins to interfere with the insert 160 to spread bullet portion 14 of the shaft to allow for clear staple release.

**90 degrees:** Former 52 fully forward; staple ejected from the device.

25

The use of cams in cam mechanism 62 ensures the accuracy of sequence and relative timing between events as well as ensuring positive mechanical displacements of all components.

30

If desired, Fig. 17, the bullet head 14 of the shaft 10, which approximates the blood vessel wall 208, may include a number of suction ports 200. These ports are

in communication with a suction adapter 202 via capillaries 204 within the shaft section. Suction, from a standard wall suction outlet or independent suction pump, is supplied to the suction adapter 202 via an on/off tap 206. Once the device is in position on the arterial wall, as indicated by blood flowing from the blood exit port, the tap 206 is turned to the "on" position thereby delivering suction to the ports 200 on the bullet head 14. This in turn suctions the blood vessel wall 208 against the face of the head 14 so as to stabilise it during delivery of the staple. Once delivered the suction is deactivated so as to remove the device from the blood vessel wall and tissue tract.

15

An alternative embodiment for the staple 40 is illustrated in Figs. 20(A) and 20(B). In this case the centre portion 40C of the base section 40A is deformed in a generally 'U' shaped loop perpendicular to the common plane of the legs 40B which form an internal angle of between 70 and 85° with the base section 40A.

Another embodiment of staple is illustrated in Figs. 21(A) and 21(B). The omega hump 40C is identical to that shown in Fig. 14. However, in this case, at the points A and B the outer ends 40A' of the base section are pre-bent backwardly to make an angle of between 130° and 150° with the inner ends 40A". The legs 40B extend forwardly from the outer ends 40A' of the base section and are substantially parallel to one another and perpendicular to a plane containing the omega section 40C and inner ends 40A" of the base section. The advantage of this configuration is that

the rearwardly angled base section offers stability to the staple as it is advanced forward toward the anvil in the track formed between the top and bottom shaft.

This prevents the legs tilting up or down as they

5 advance beyond the free end of the shaft. In addition, once the staple is formed, Fig. 21B, the legs 40B are angled rearwardly toward the omega hump and away from the internal lumen of the vessel.

10 Another embodiment of staple is illustrated in Figs. 22(A) and 22(B). The omega hump 40C is identical to that shown in Fig. 14. In this case, however, the inner ends 40A" of the base section are inclined back at between 30° and 50° from the plane of the omega  
15 section 40C. The base section includes the same pre-bends at points A and B between the base section inner and outer ends 40A", 40A' as shown in Fig. 14. The legs 40B extend forwardly from the outer ends 40A' of the base section and are substantially parallel to one  
20 another and perpendicular to a plane containing the omega section 40C. This construction offers the advantages of stability as the staple is advanced forward during firing, maximised closure from the pre-bend and, in the closed staple, legs 40B which are  
25 angled rearwardly toward the omega hump and away from the internal lumen of the vessel.

Yet another embodiment of staple is shown in

Figs. 23(A) and 23(B). In this case the outer ends

30 40A' of the base section and the respective legs 40B are combined to form one continuously curved element. The curved section begins at the point where the wire is intended to bend around the anvil and terminates at

a pointed free end. The advantage of this curved base/legs is they again offer stability to the staple as it is advanced forward toward the anvil in that the pointed ends advance in a plane perpendicular to the omega section. Also, because of the curved nature of the base/leg the pointed end is always tending to advance into and across the puncture hole in the wall as opposed to advance through the wall and into the lumen and finally, after forming the leg points are facing away from the lumen and toward the hump section.

The embodiments shown herein have a loop which is substantially perpendicular to the common plane, although this need not be the case. The loop is preferably at an angle of at least 75 degrees from the common plane, as this enables it to lie close to e.g. an arterial wall when applied. For this reason a substantially perpendicular loop is preferred, but for design reasons a loop lying at any other angle could be employed provided that the staple can be properly deployed and that the loop does not cause unnecessary trauma when applied.

Obviously, features such as the bruise and pre-bend points can be applied to any of the embodiments shown above. While the illustrations show staples fabricated from round section wire the invention is not restricted to this. Other profiles such as rectangular, square, triangular, etc can be employed.

30

Furthermore, the staple need not be symmetrical about the loop, i.e. the base section and/or the legs can be different on the left-hand side and right-hand side.

This may be particularly desirable where the staple is applied at an angle to the tissue wall and thus each leg is required to travel a different distance or to be deformed in a different shape to accomplish an optimal  
5 closure.

The invention is not limited to the embodiments described herein and may be modified or varied without departing from the scope of the invention.

**Claims**

1. A surgical staple comprising a base portion and a  
respective leg extending from each end of the base  
5 portion and terminating at a free end, the base portion  
being deformable to bring the free ends of the legs  
together to penetrate a liquid-carrying vessel at the  
site of a puncture and hold the opposite edges of the  
puncture together, the base portion and legs lying in  
10 substantially a common plane except for a centre  
portion of the base portion which is deformed in a loop  
at an angle to the common plane.
2. A surgical staple as claimed in claim 1, wherein  
15 the loop is generally U-shaped.
3. A surgical staple as claimed in claim 1, wherein  
the loop is generally omega-shaped.
- 20 4. A surgical staple according to claim 1, 2 or 3,  
wherein each leg extends at an angle from the  
respective end of the base portion, the legs pointing  
at least approximately in the same direction.
- 25 5. A surgical staple as claimed in claim 4, wherein  
the legs are substantially parallel.
6. A surgical staple as claimed in claim 4, wherein  
the legs converge at a small angle.
- 30 7. A surgical staple as claimed in claim 4, 5 or 6,  
wherein on each side of the loop the outer end of the

base portion is pre-bent at a small angle relative to the inner end of the base portion.

8. A surgical staple as claimed in claim 7, wherein  
5 the outer portion is bent in the opposite direction to the direction in which the legs point.

9. A surgical staple as claimed in claim 7, wherein  
10 the outer portion is bent in the same direction as the direction in which the legs point.

10. A surgical staple as claimed in any preceding  
claim, wherein the loop lies in a plane at an angle of  
at least 75 degrees to the common plane.

15 11. A surgical staple as claimed in any preceding  
claim, wherein the loop lies in a plane substantially  
perpendicular to the common plane.

20 12. A surgical staple as claimed in any preceding  
claim, wherein the legs form a continuous curve with  
the outer end of the base portion.

13. A surgical staple as claimed in any preceding  
25 claim, wherein on each side of the loop the base  
portion is deformed to encourage bending at that point  
during closure of the staple.

14. A method of stapling closed a puncture site in a  
30 liquid-carrying vessel in a human or animal body,  
comprising the steps of:

introducing a stapling mechanism to the location  
of the vessel;

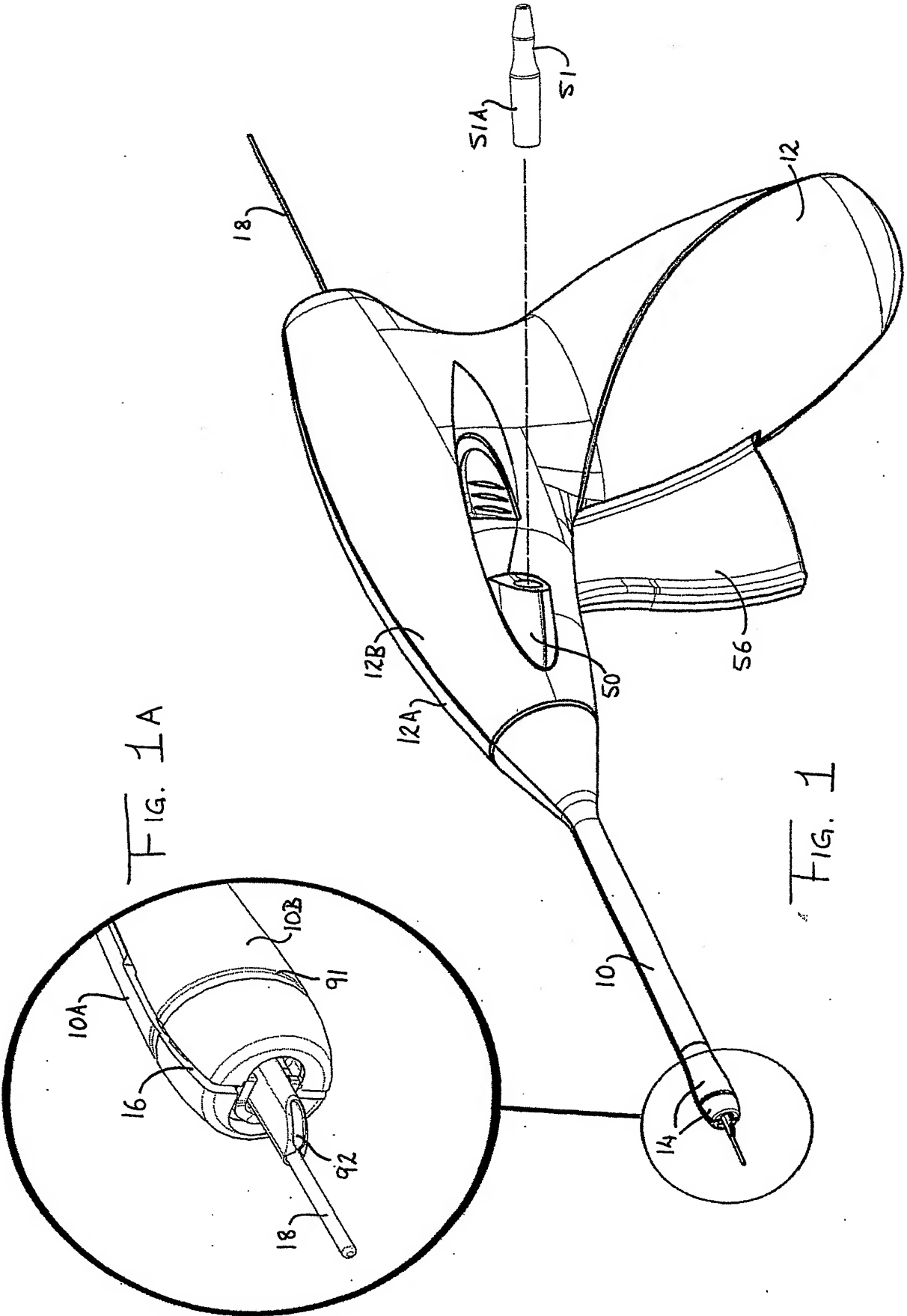
positioning the stapling mechanism at the puncture site by means of an elongated locator device associated with the stapling mechanism, the locator device sensing the position of the puncture site by entering the

5 vessel at the site; and

delivering a staple to, and deforming the staple to close, the puncture site;

wherein the staple comprises a base portion and a respective leg extending from each end of the base  
10 portion and terminating at a free end, the base portion being deformable to bring the free ends of the legs together to penetrate a blood vessel at the site of a puncture and hold the opposite edges of the puncture together, the base portion and legs lying in  
15 substantially a common plane except for a centre portion of the base portion which is deformed in a loop at an angle to the common plane; and

wherein during delivery of the staple to the puncture site the staple loop straddles and slides  
20 along the locator device.



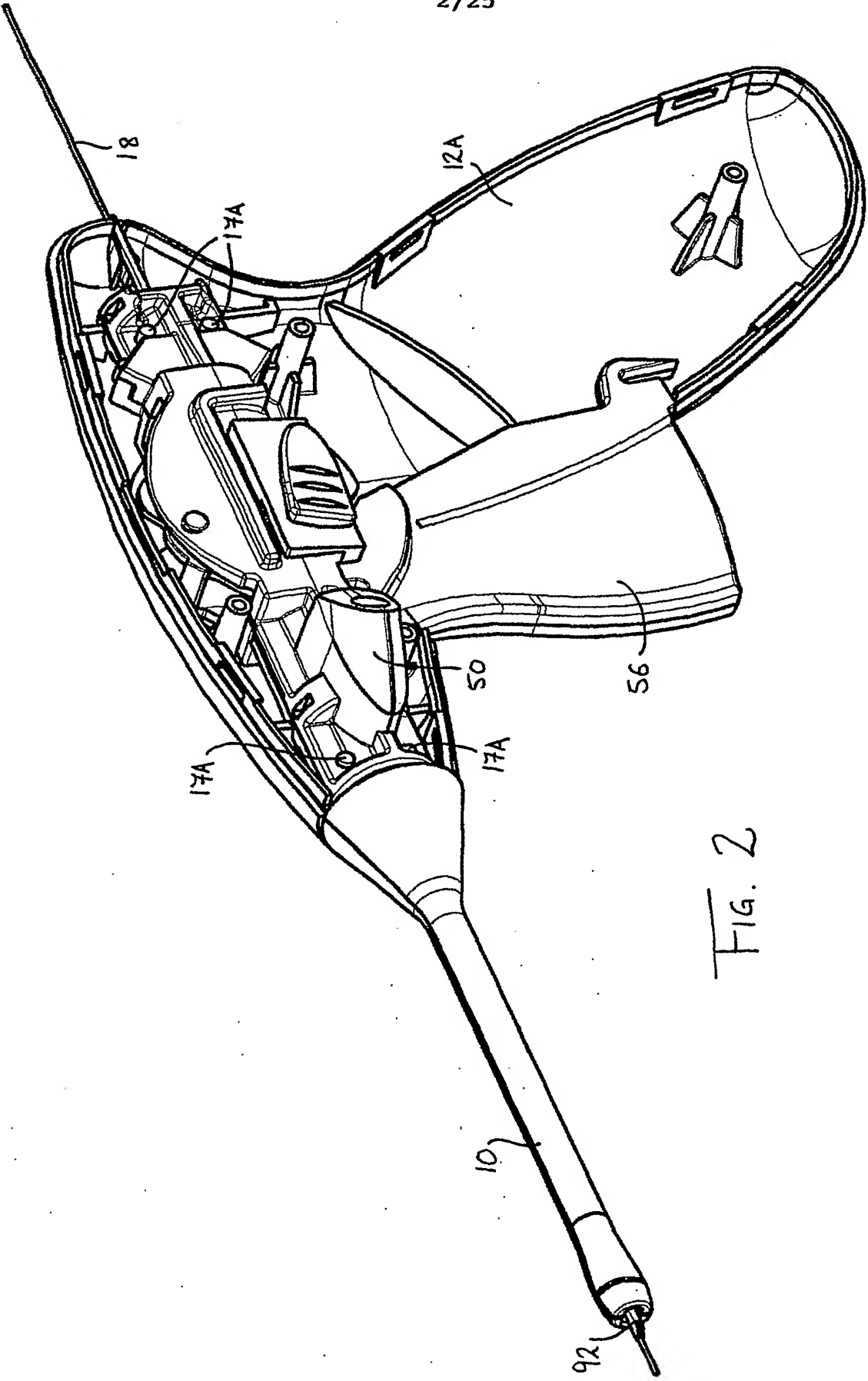
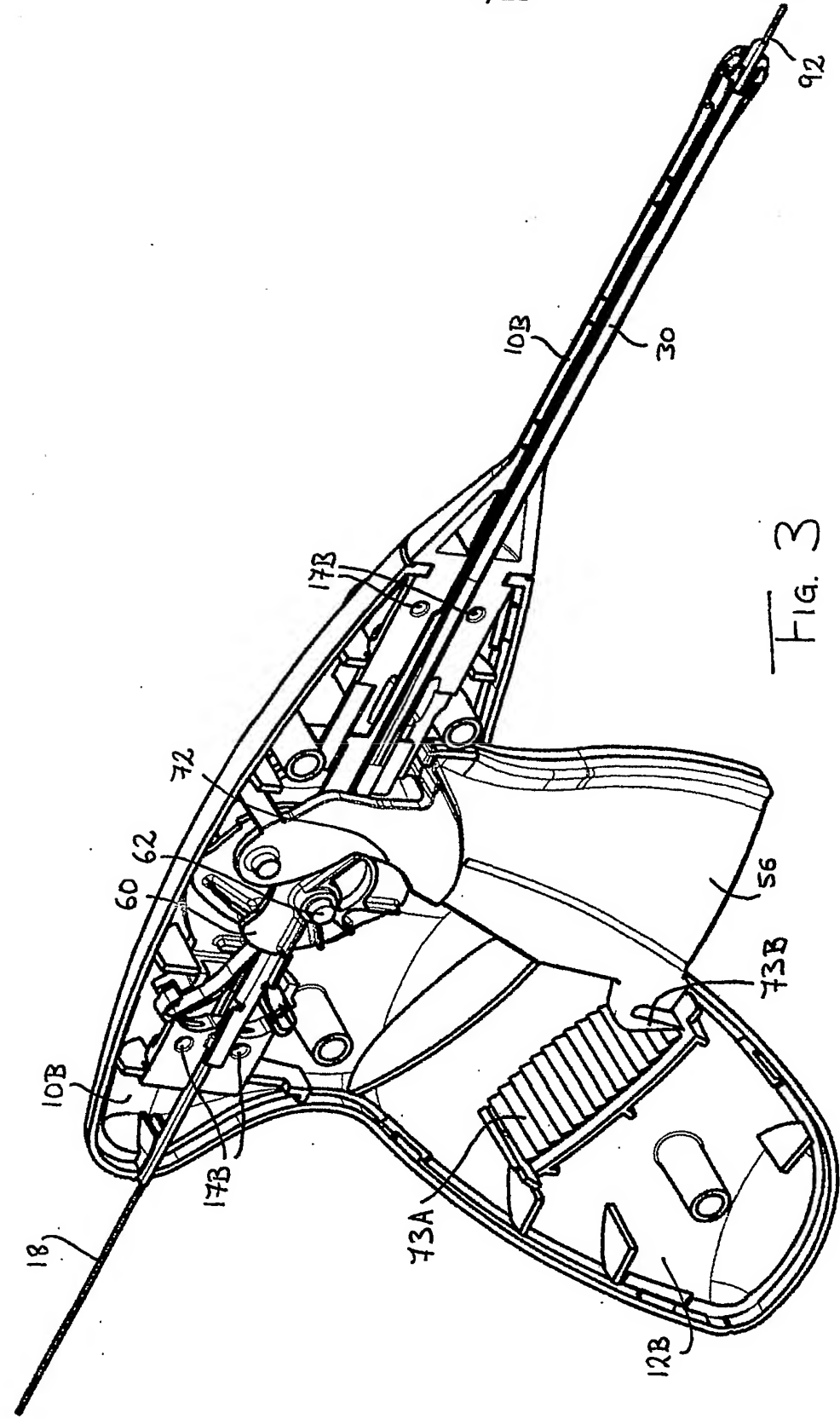
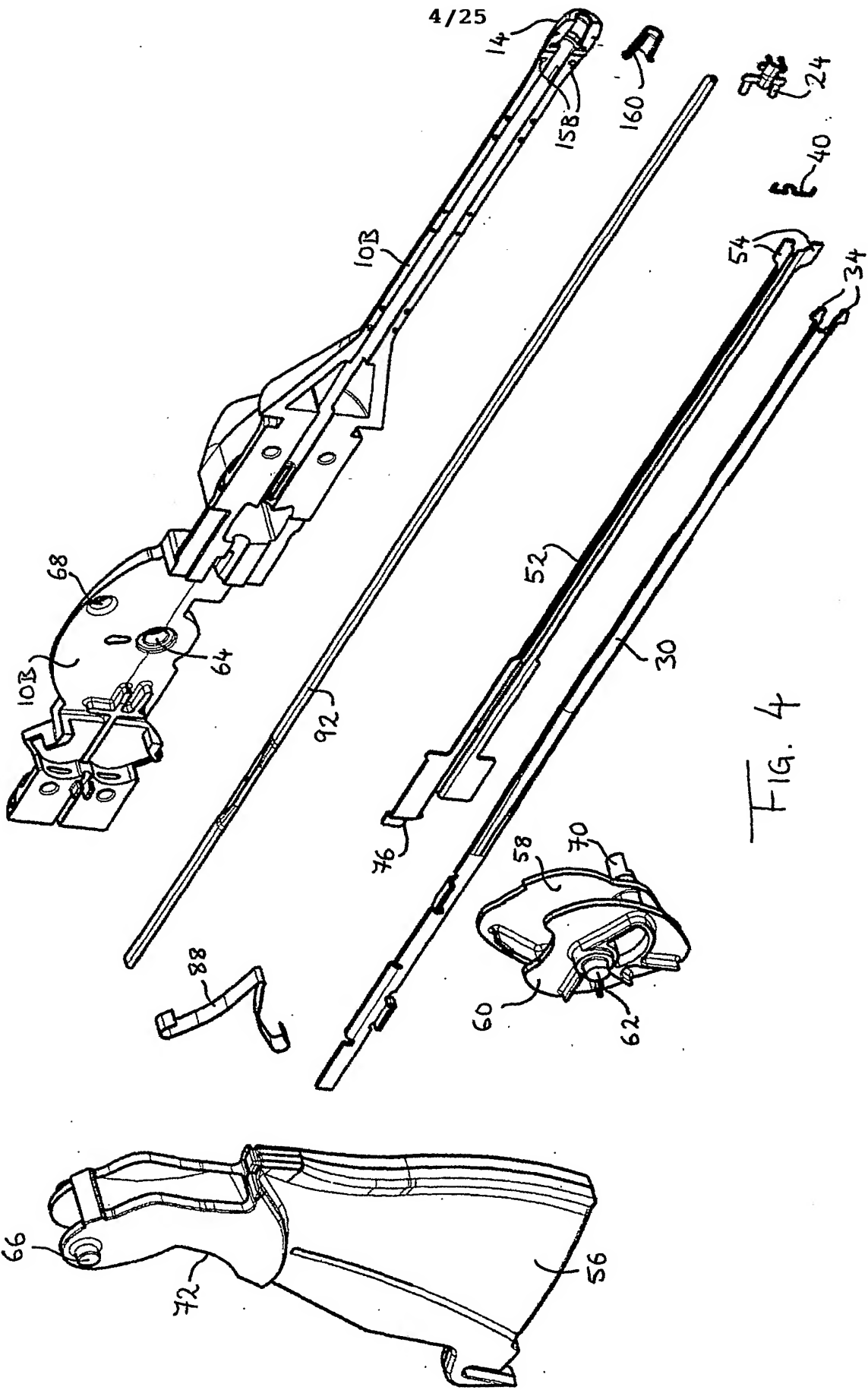


FIG. 2





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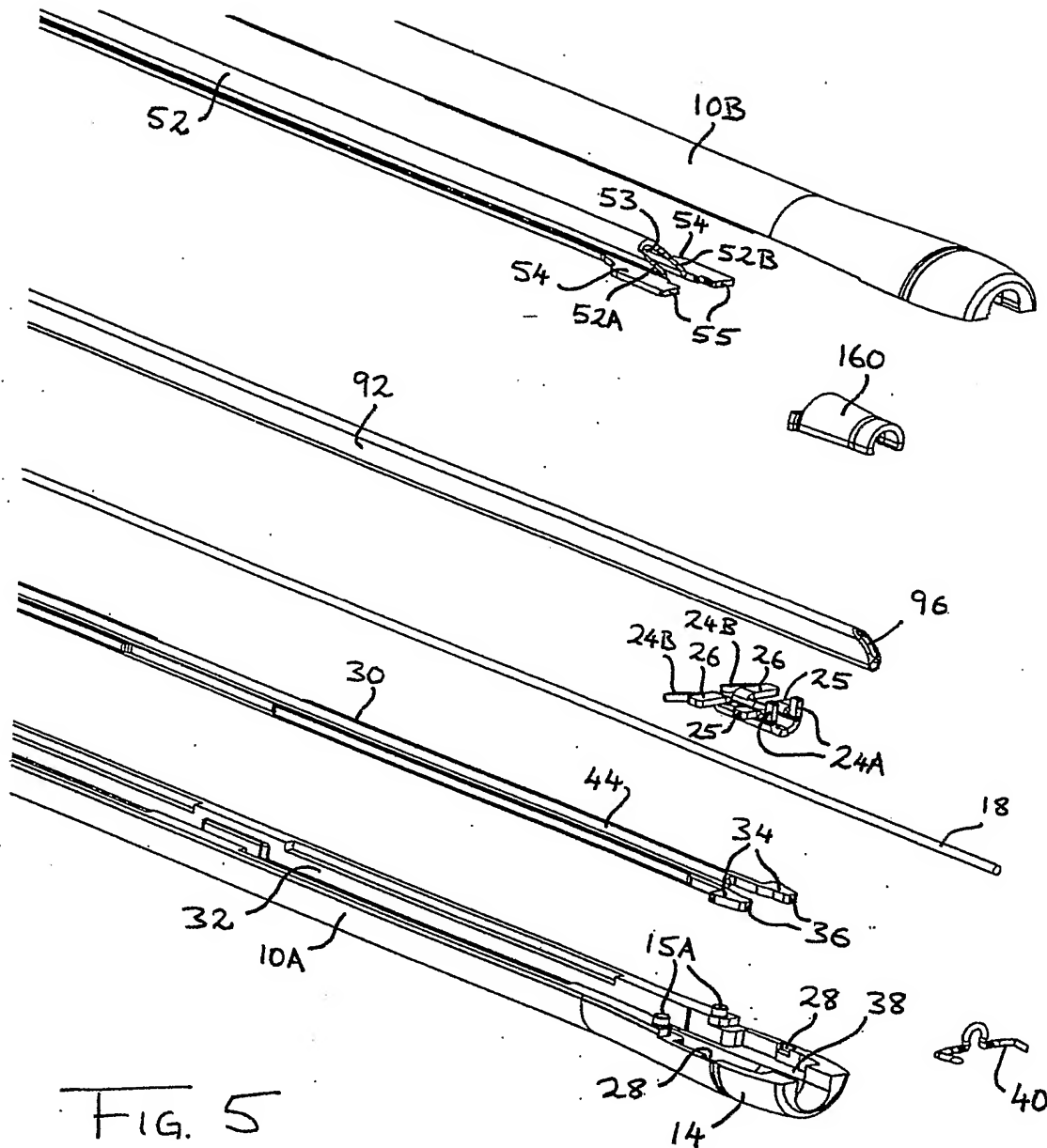
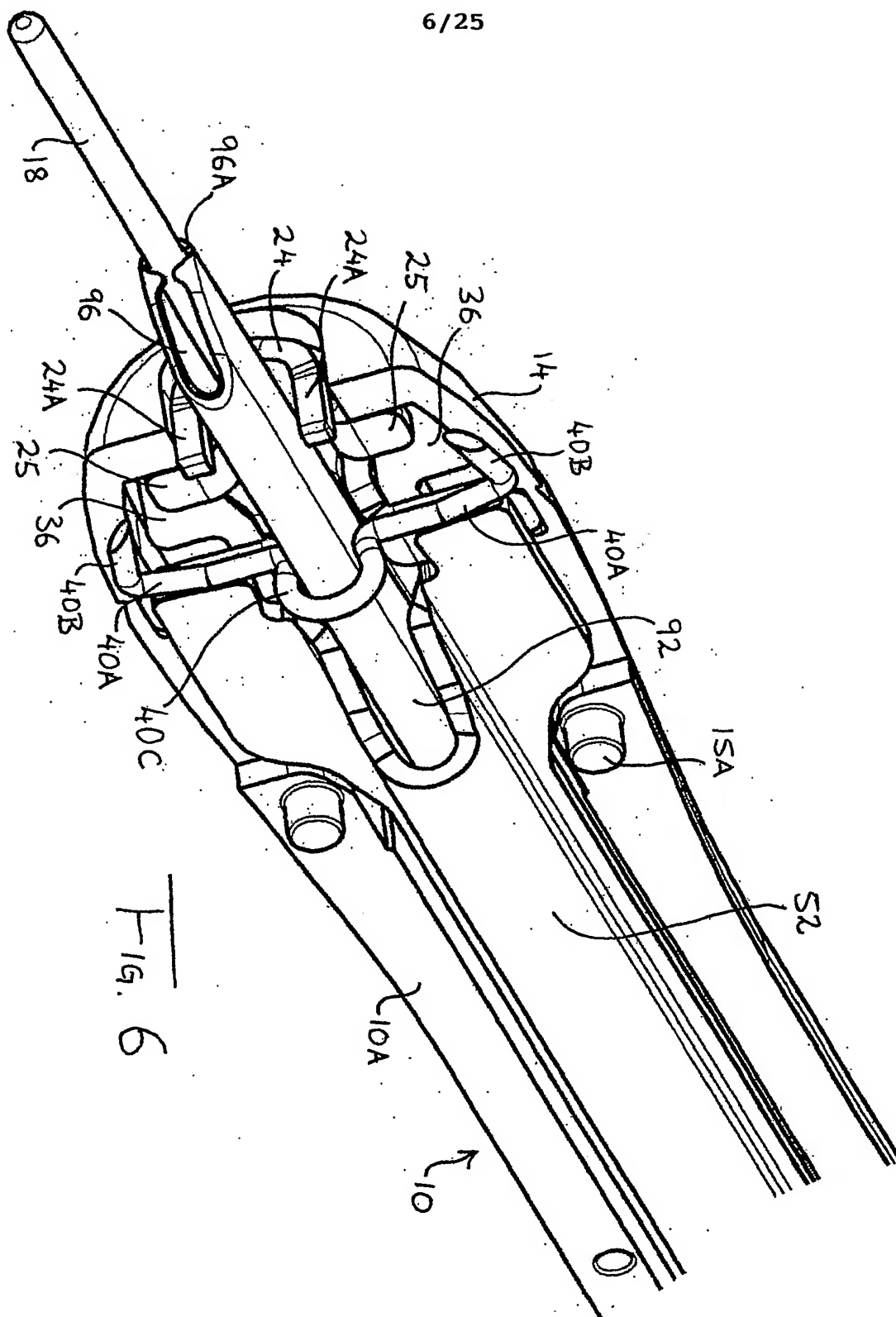


FIG. 5

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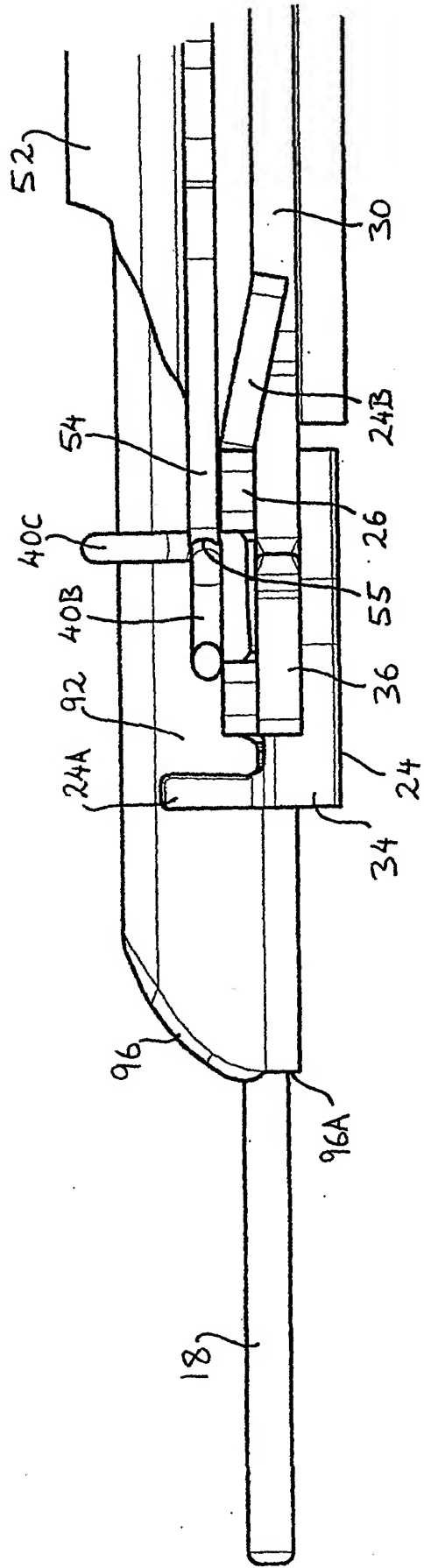
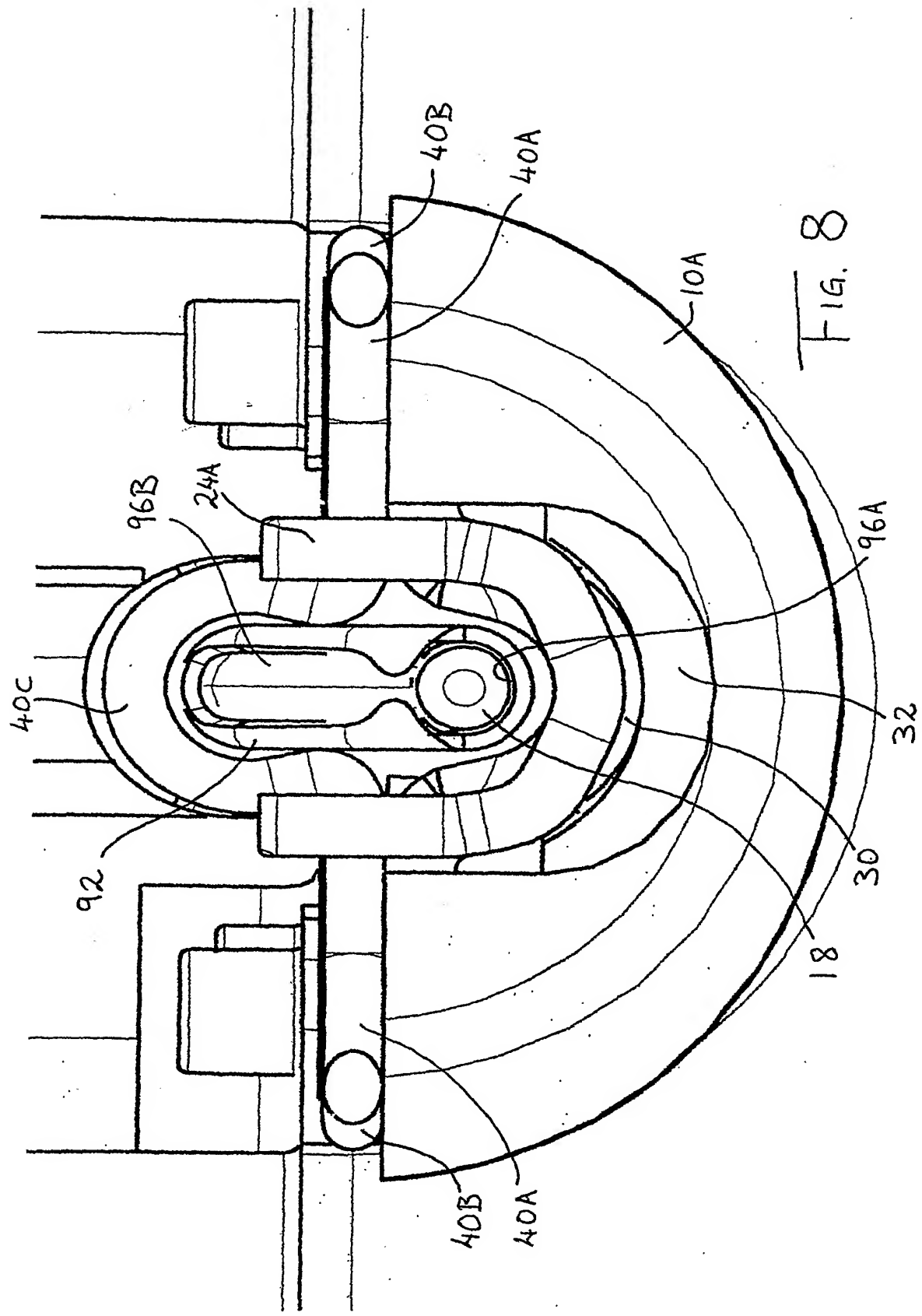


FIG. 7



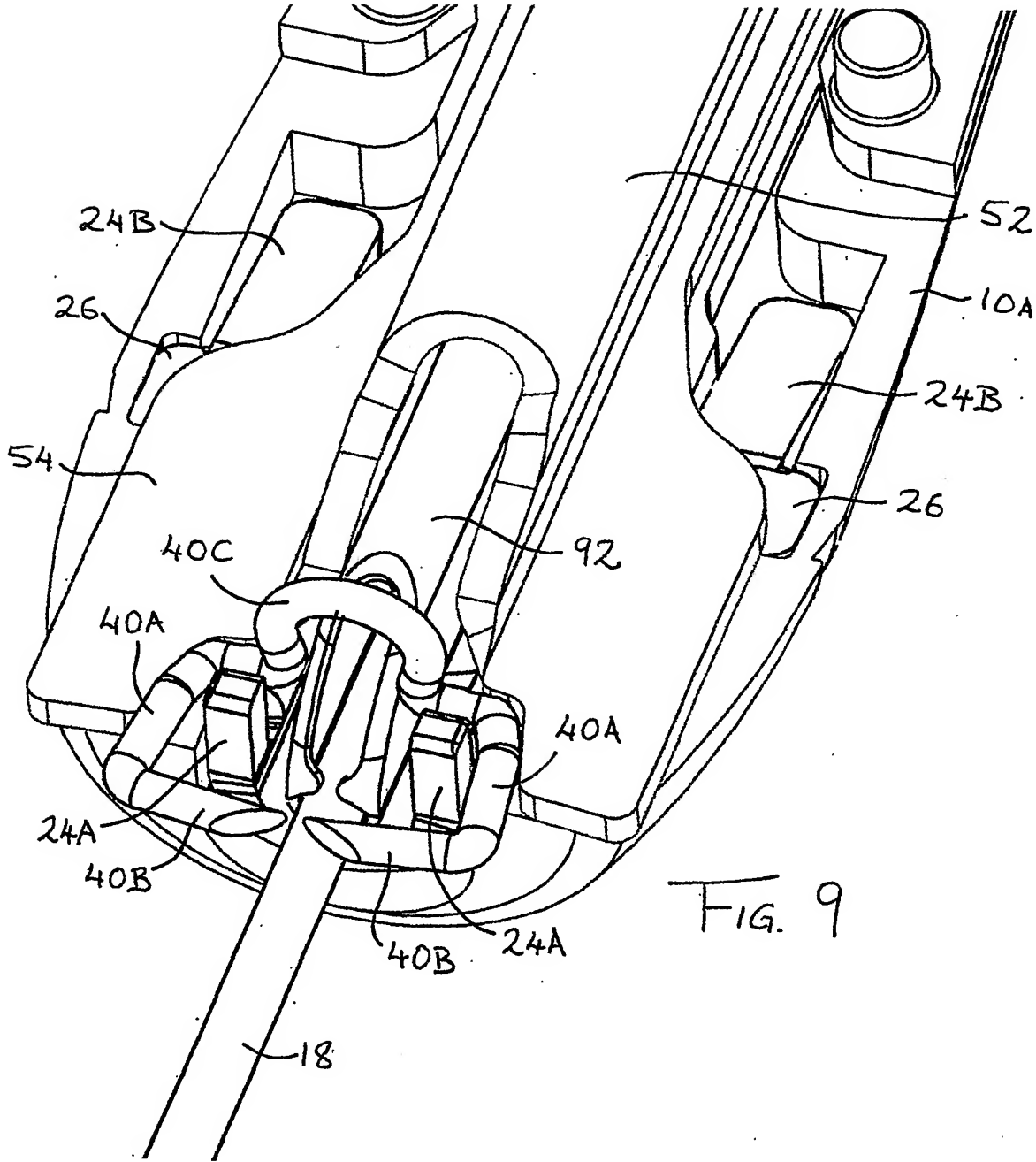
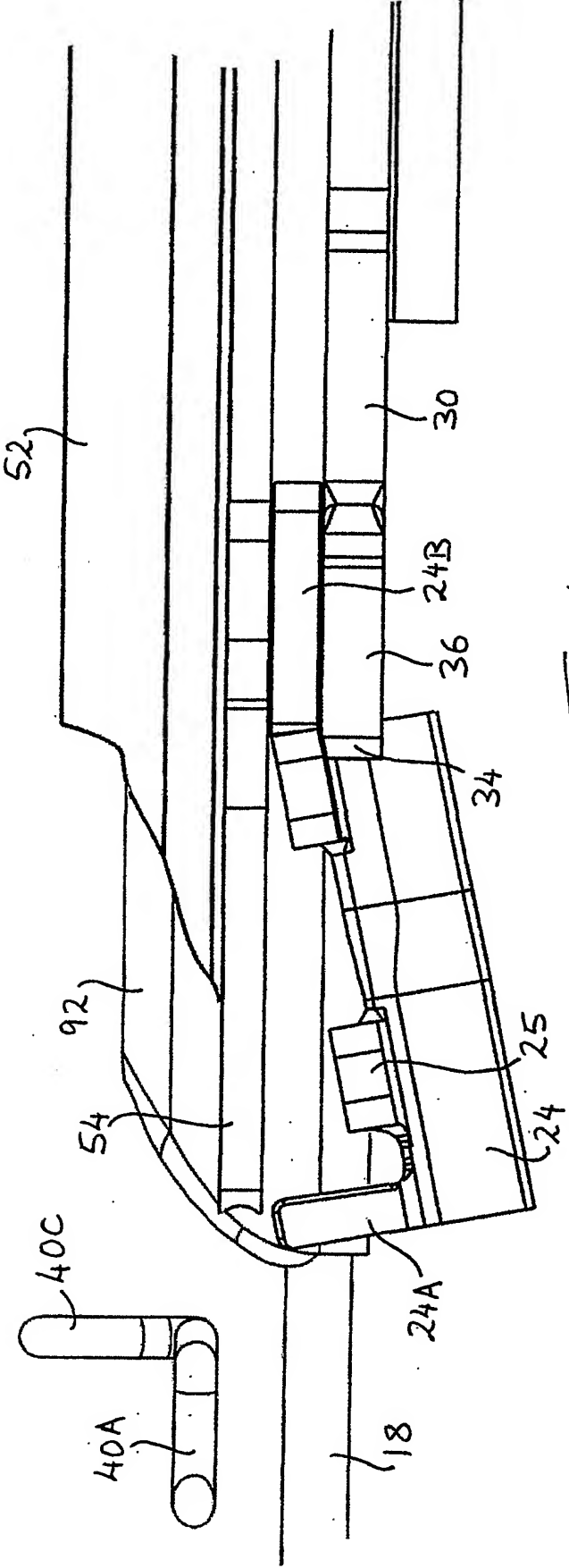
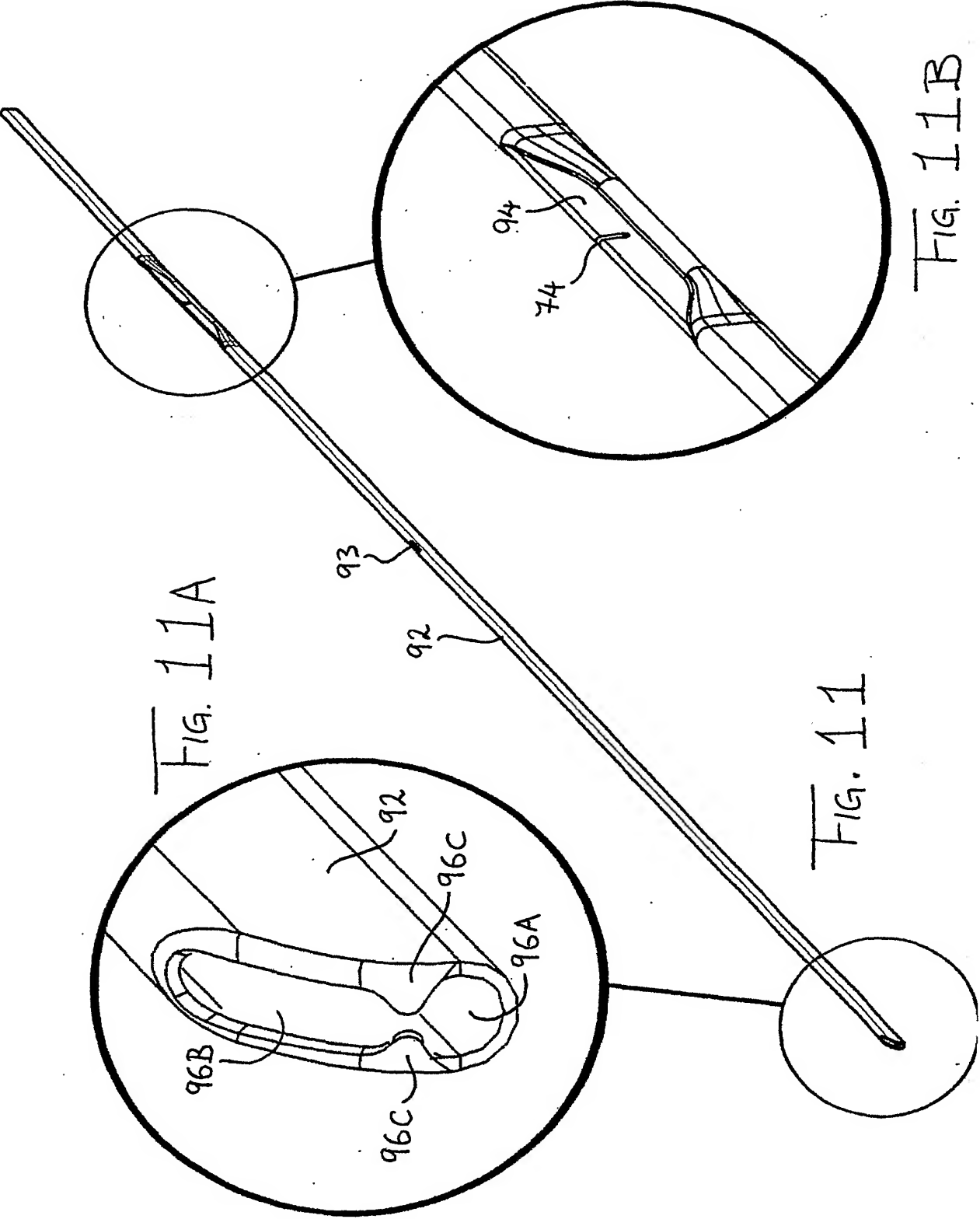


FIG. 9



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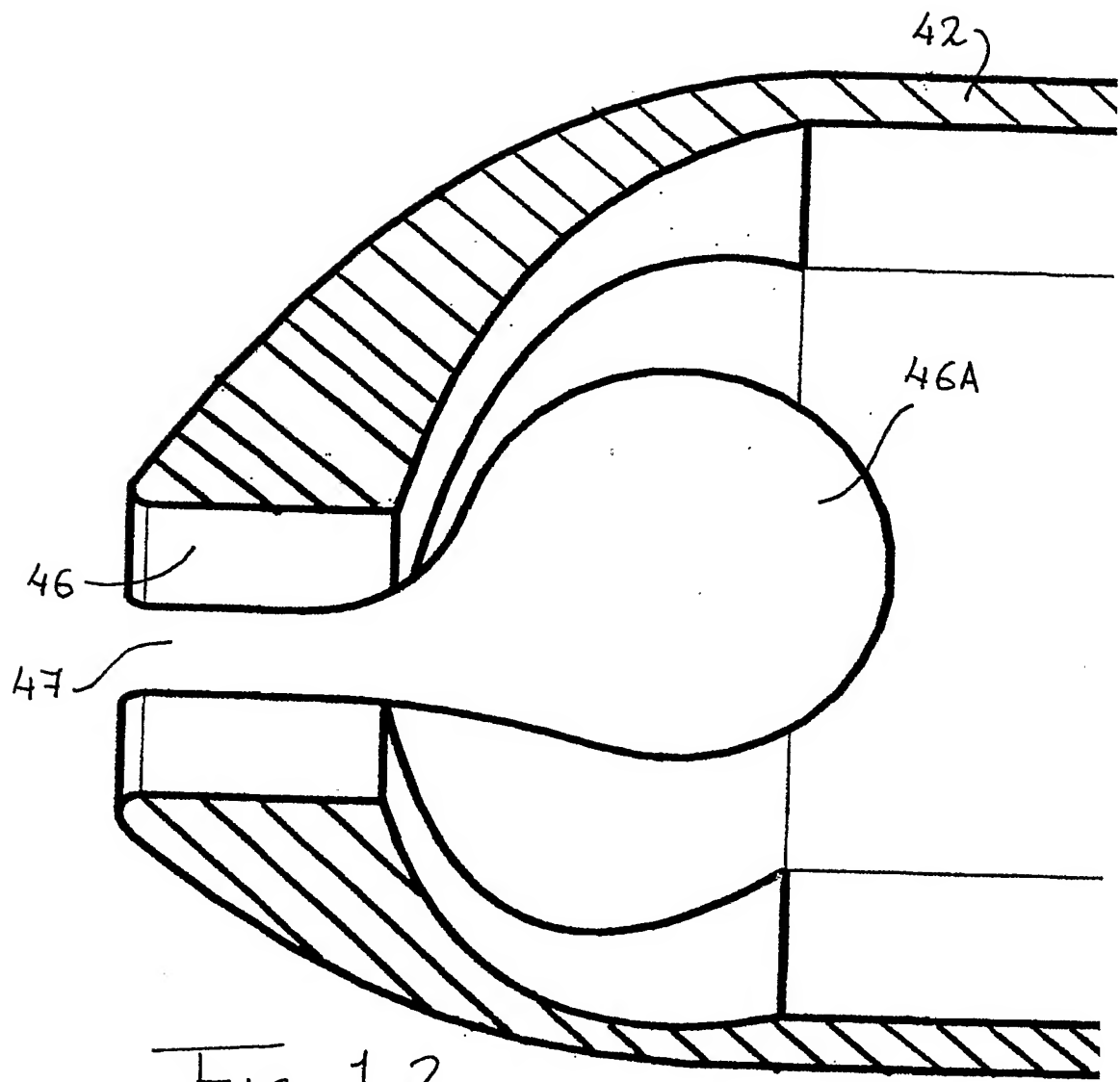


FIG. 12

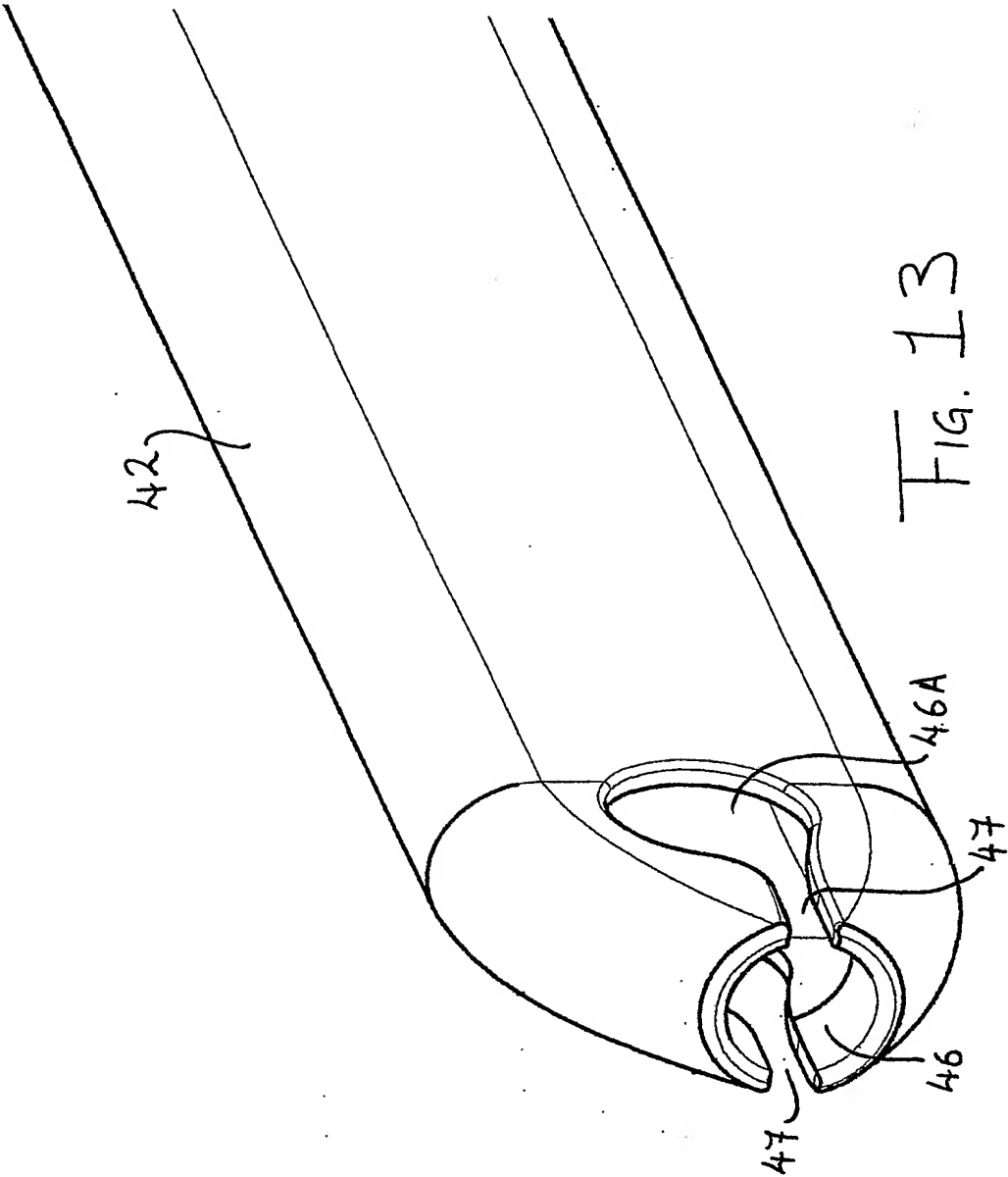
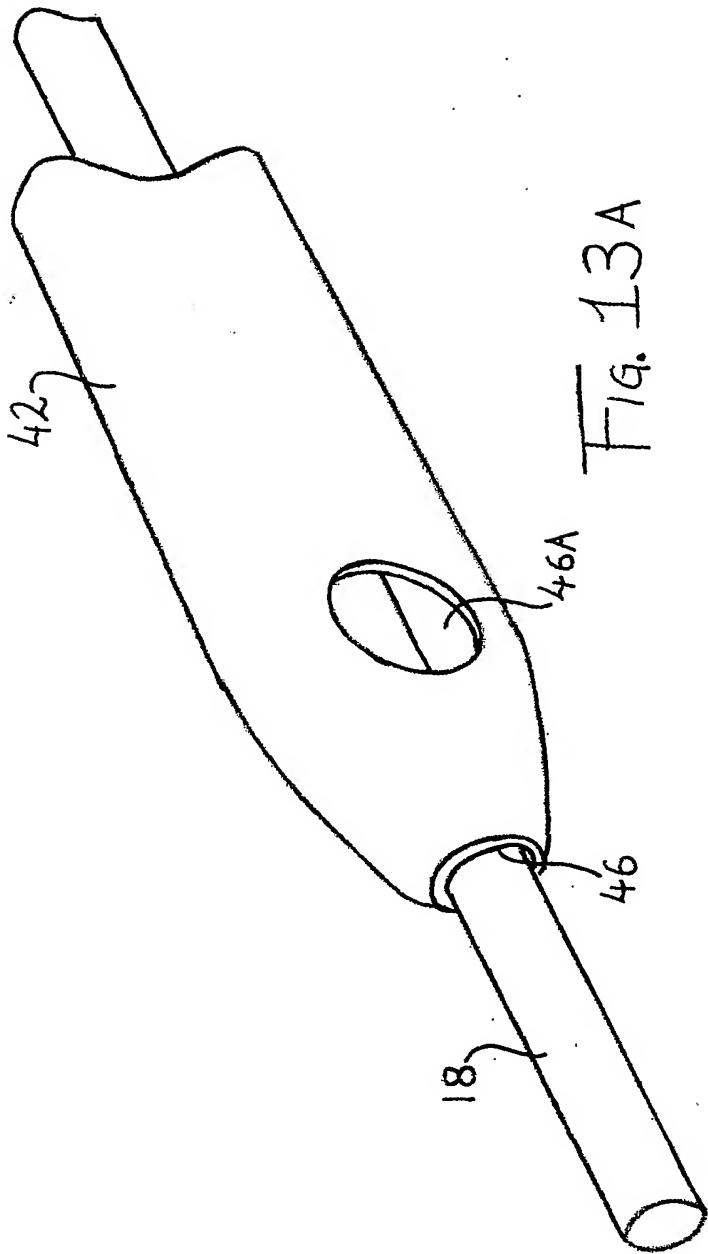


FIG. 13



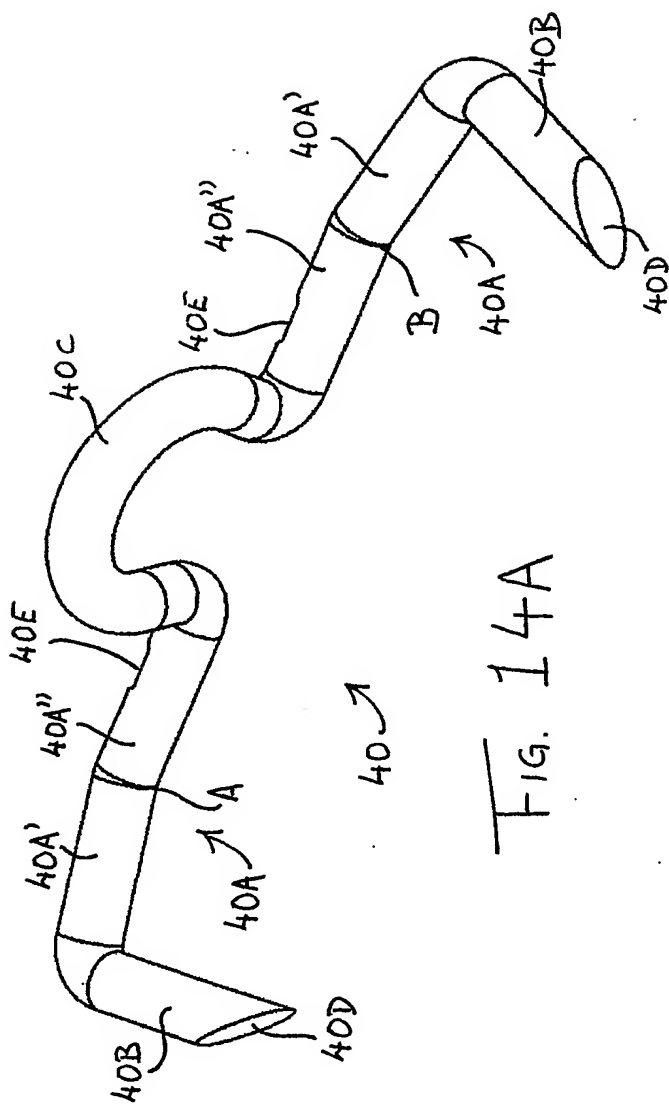


FIG. 14A

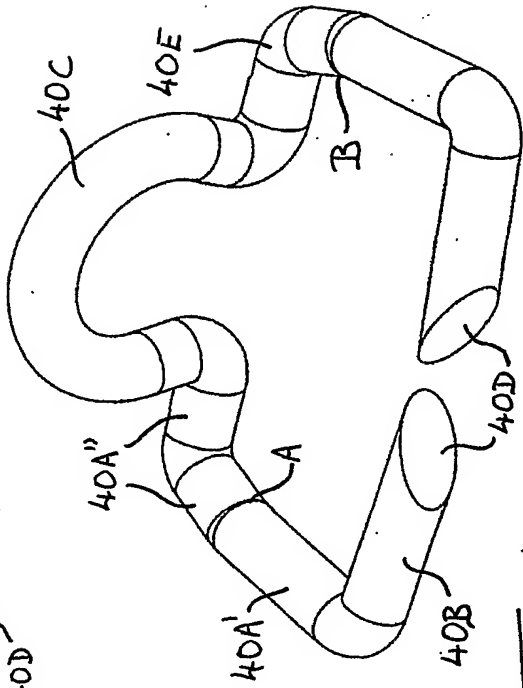
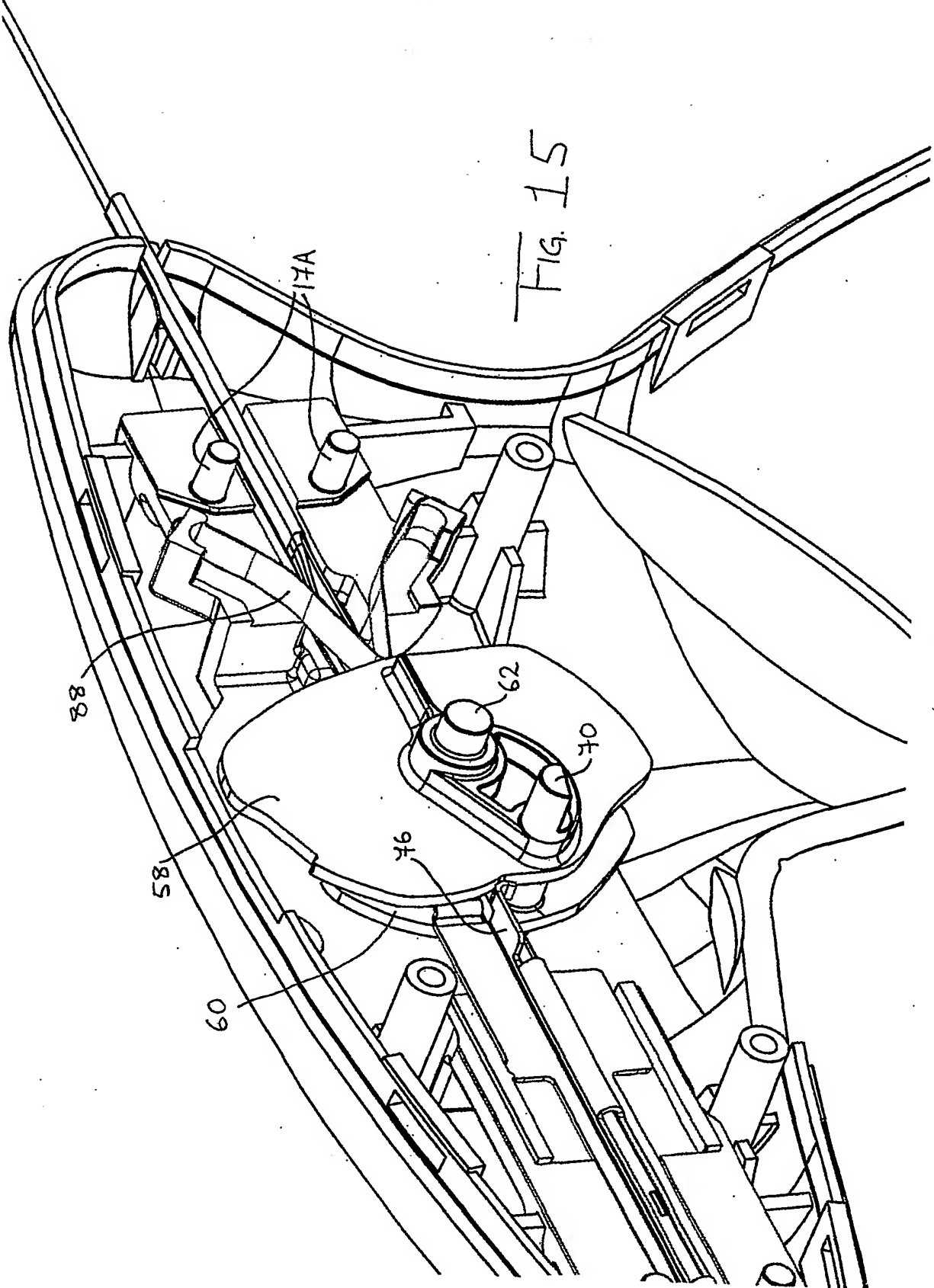


FIG. 14B

FIG. 15



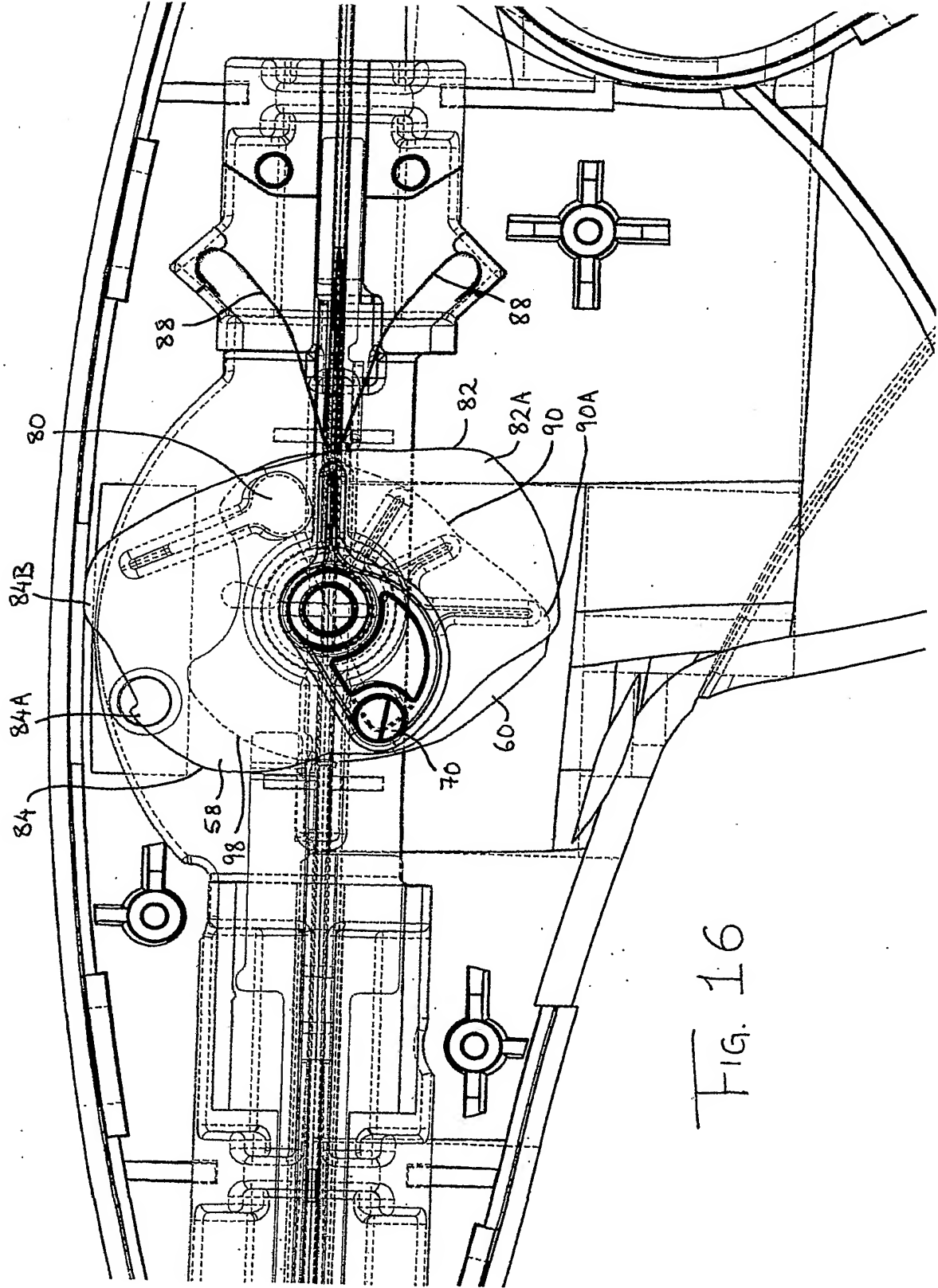
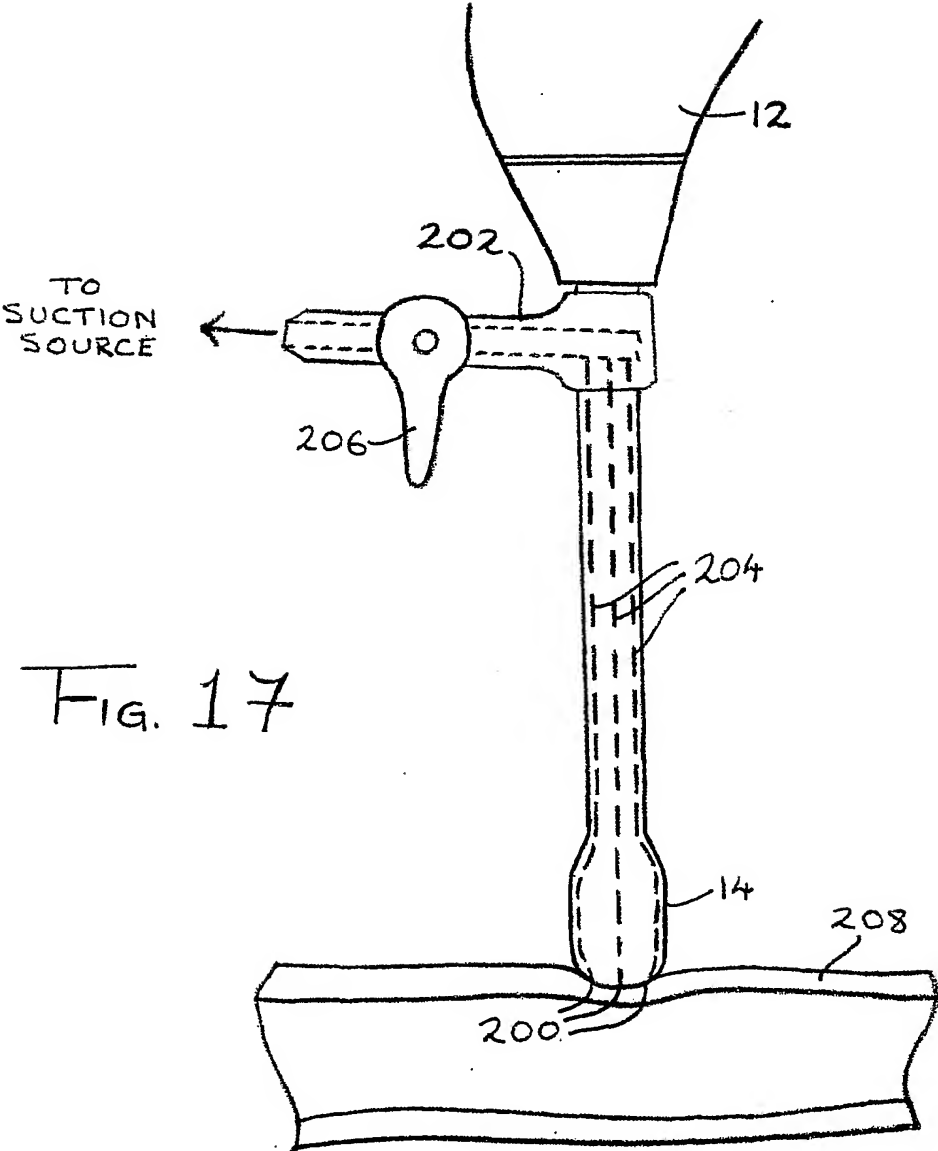


FIG. 16



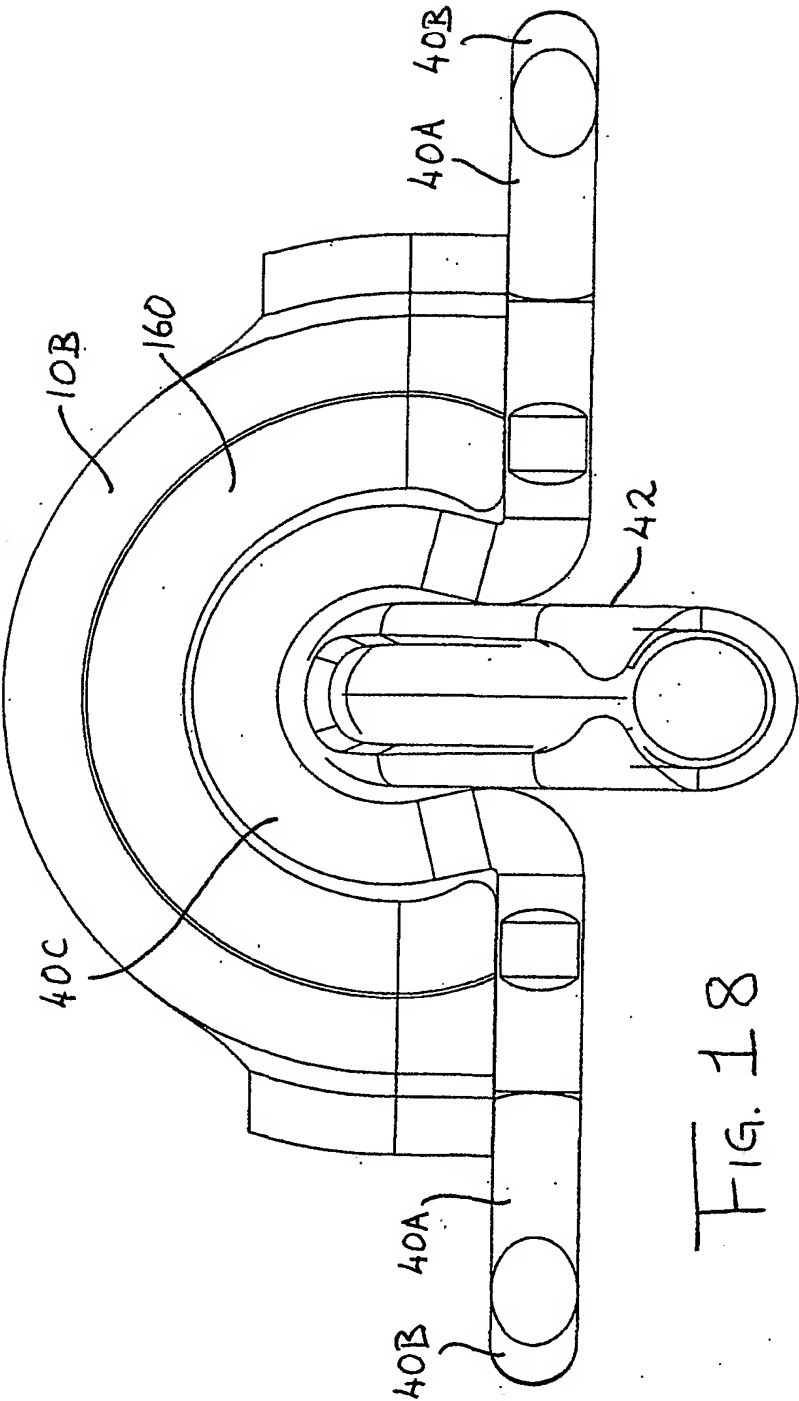


FIG. 18

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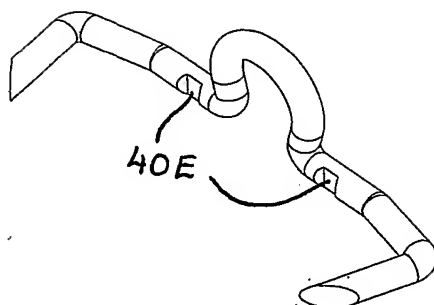


FIG. 19A

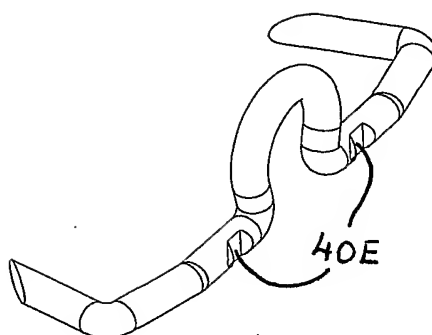


FIG. 19B

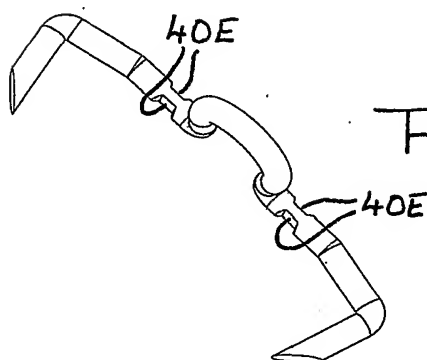


FIG. 19C

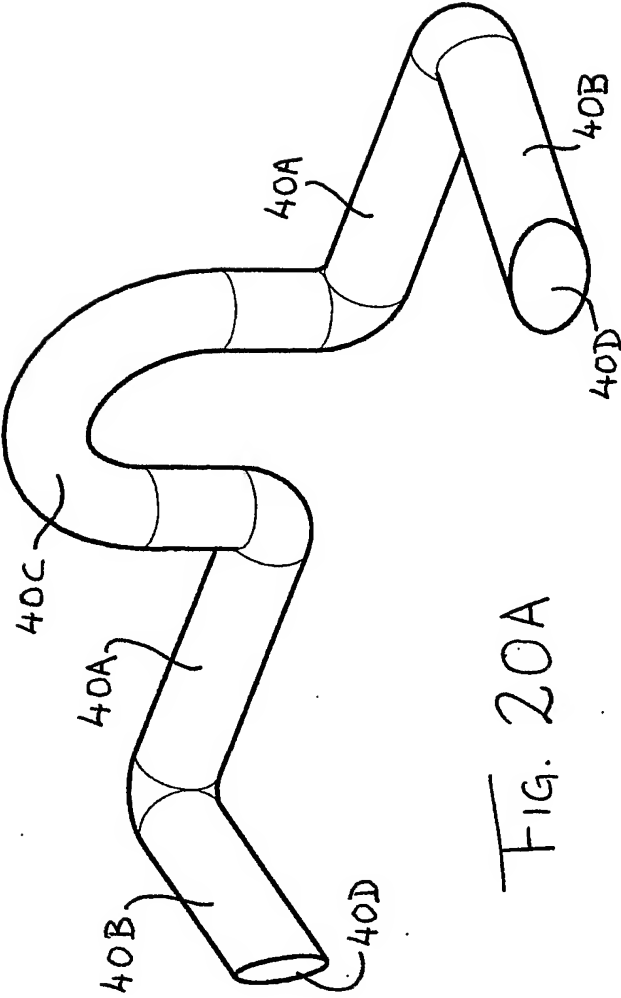


FIG. 20A

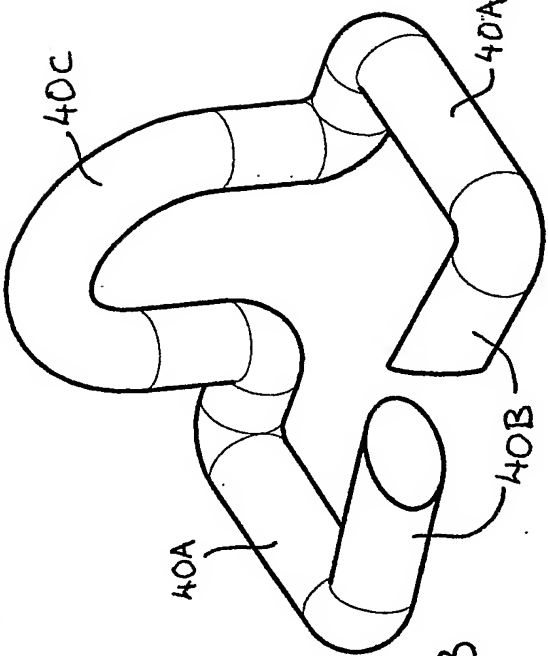
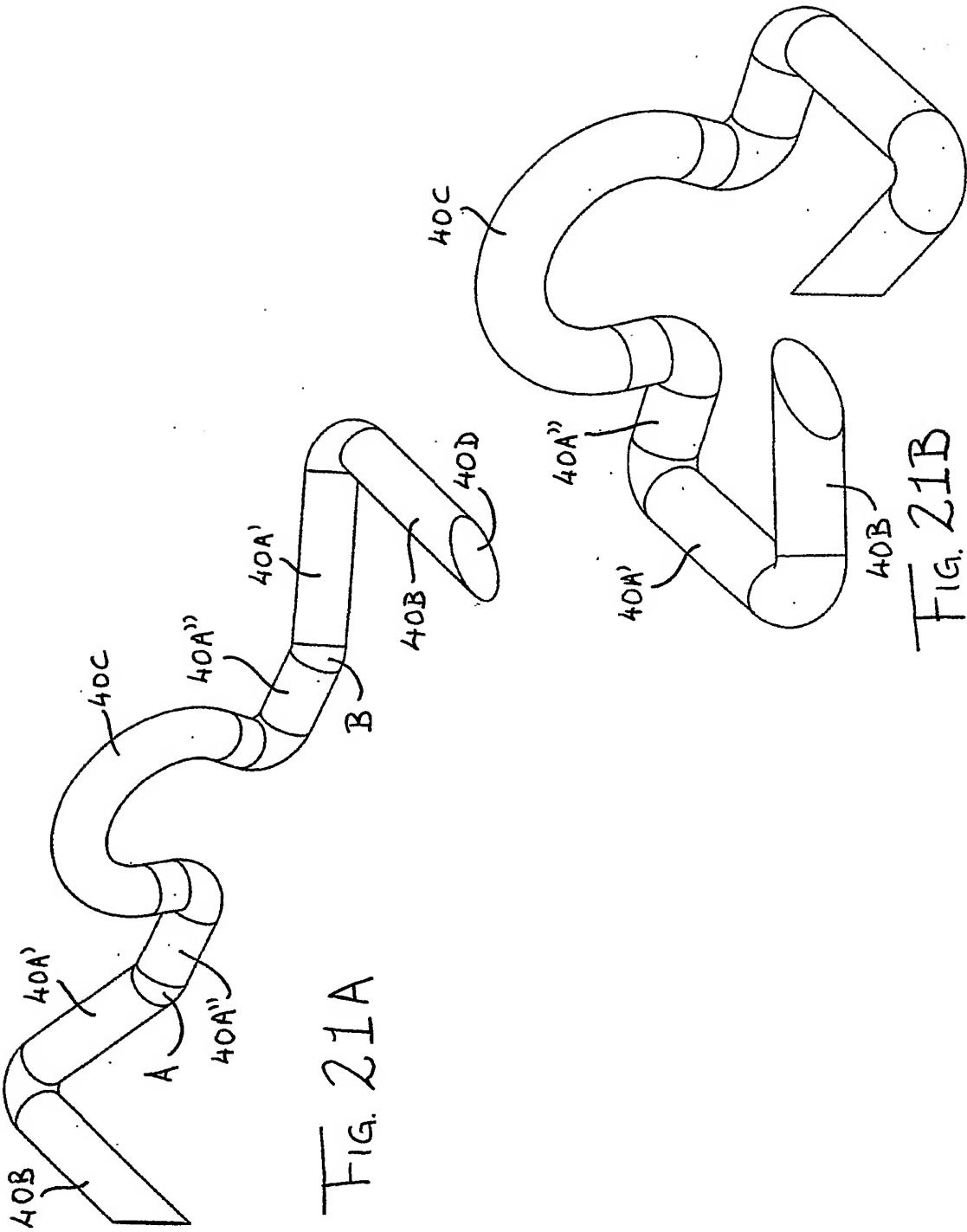
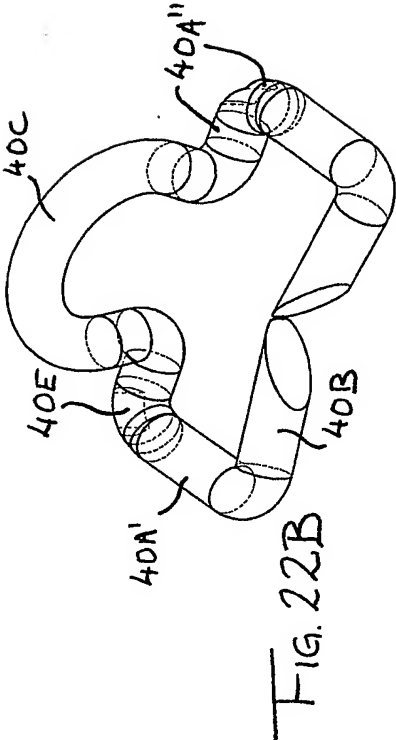
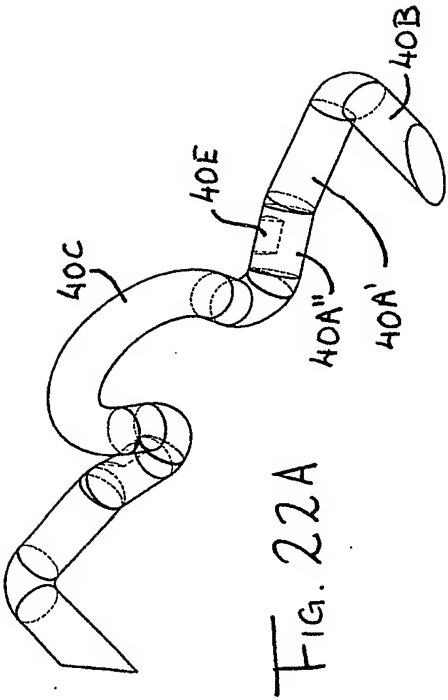


FIG. 20B





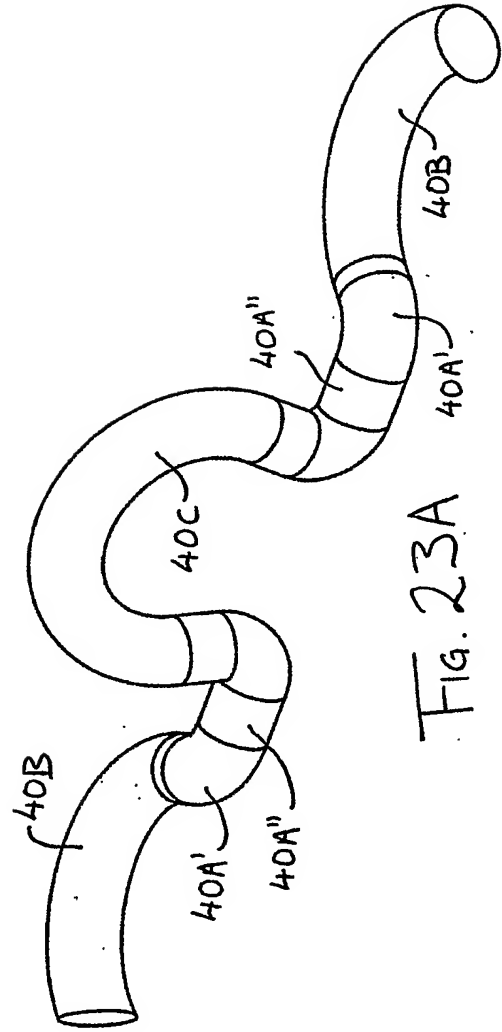


FIG. 23A

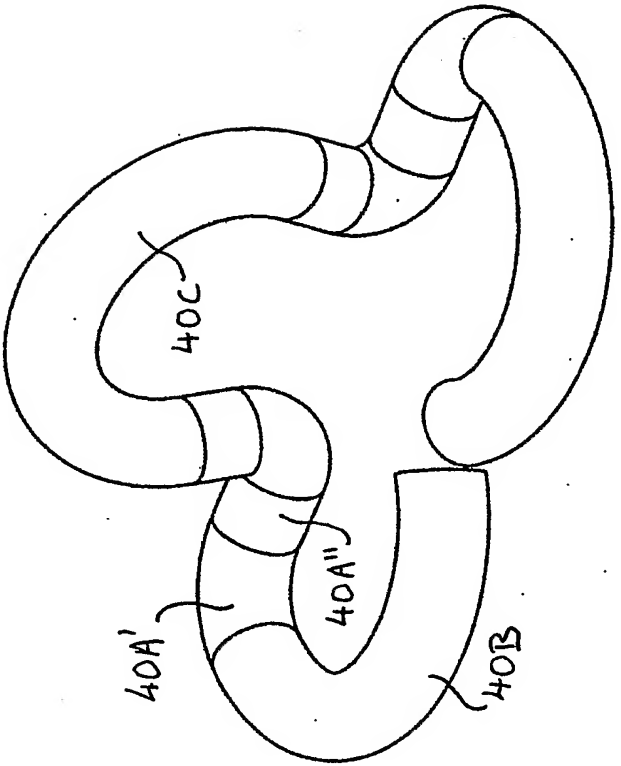


FIG. 23B

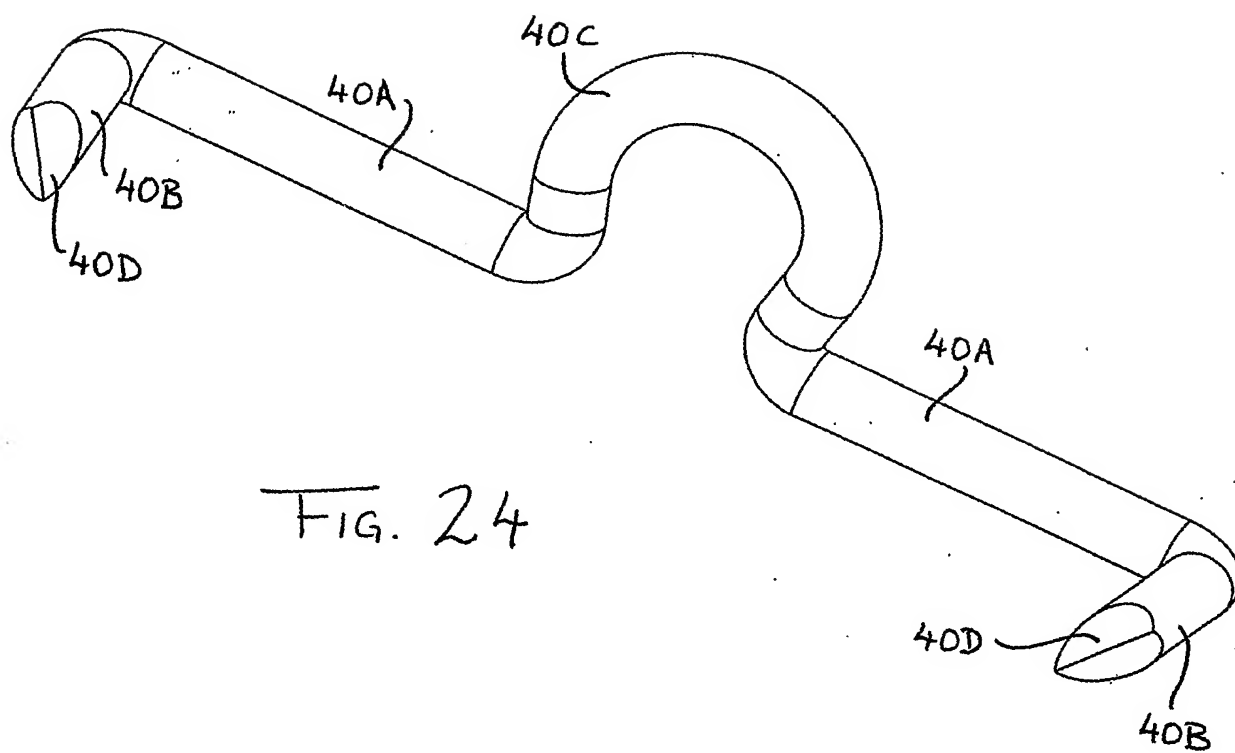


FIG. 24

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/IE 01/00117

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61B17/064 A61B17/068 A61B17/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 386 361 A (ETHICON INC) 12 September 1990 (1990-09-12)	1-6, 11, 12
Y	the whole document	7-9, 13
Y	US 5 470 010 A (ROTHFUSS ROBERT G ET AL) 28 November 1995 (1995-11-28)	7, 9, 13
A	abstract; figure 13	1
Y	US 4 014 492 A (ROTHFUSS ROBERT G) 29 March 1977 (1977-03-29)	8
A	abstract; figure 1	1
A	US 5 147 381 A (HEIMERL ALBERT ET AL) 15 September 1992 (1992-09-15)	1, 6
	abstract; figure 2	
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☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

15 November 2001

Date of mailing of the international search report

23/11/2001

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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/IE 01/00117

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98 25508 A (ALLGEYER DEAN OWEN) 18 June 1998 (1998-06-18) abstract; figures 1,2 ---	1
A	US 5 560 532 A (DEFONZO STEPHAN A ET AL) 1 October 1996 (1996-10-01) abstract; figures 13-16 -----	1

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IE 01/00117

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0386361	A	12-09-1990	CA 2000360 A1	11-04-1990
			EP 0386361 A1	12-09-1990
			GR 89100652 A	29-11-1990
			IE 327089 L	11-04-1990
			PT 91958 A	30-04-1990
<hr/>				
US 5470010	A	28-11-1995	US 5333772 A	02-08-1994
			US 5246156 A	21-09-1993
			US 5174487 A	29-12-1992
			AT 179317 T	15-05-1999
			AU 652276 B2	18-08-1994
			AU 2213392 A	25-03-1993
			AU 672535 B2	03-10-1996
			AU 7897794 A	02-02-1995
			CA 2077964 A1	13-03-1993
			DE 69229031 D1	02-06-1999
			DE 69229031 T2	23-09-1999
			EG 19763 A	29-02-1996
			EP 0536903 A2	14-04-1993
			ES 2130159 T3	01-07-1999
			GR 92100363 A , B	21-06-1993
			JP 5192340 A	03-08-1993
			TR 27512 A	07-06-1995
			AT 139893 T	15-07-1996
			AU 650959 B2	07-07-1994
			AU 1397492 A	08-10-1992
			BR 9201193 A	01-12-1992
			CA 2064503 A1	05-10-1992
			DE 69211882 D1	08-08-1996
			DE 69211882 T2	28-11-1996
			EP 0507605 A1	07-10-1992
			ES 2089397 T3	01-10-1996
			GR 92100110 A , B	16-03-1993
			JP 3199831 B2	20-08-2001
			JP 5115486 A	14-05-1993
<hr/>				
US 4014492	A	29-03-1977	AR 207823 A1	29-10-1976
			AT 352252 B	10-09-1979
			AT 422776 A	15-02-1979
			AU 1467376 A	15-12-1977
			BE 842813 A1	01-10-1976
			BR 7603740 A	08-02-1977
			CA 1053110 A1	24-04-1979
			CH 610514 A5	30-04-1979
			CS 209872 B2	31-12-1981
			DD 125325 A5	13-04-1977
			DE 2625991 A1	16-12-1976
			ES 448794 A1	16-07-1977
			FR 2313903 A1	07-01-1977
			GB 1511396 A	17-05-1978
			HK 29182 A	09-07-1982
			IT 1062326 B	10-10-1984
			JP 949034 C	20-04-1979
			JP 52011682 A	28-01-1977
			JP 53027593 B	09-08-1978
			NL 7606270 A , B,	14-12-1976
			SE 414864 B	25-08-1980
			SE 7606573 A	12-12-1975

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IE 01/00117

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 4014492	A		YU 144276 A1	21-01-1983
			ZA 7603367 A	25-05-1977
US 5147381	A	15-09-1992	DE 4014653 A1	14-11-1991
			AT 111709 T	15-10-1994
			AU 626145 B2	23-07-1992
			AU 7293791 A	14-11-1991
			DE 59102997 D1	27-10-1994
			EP 0456964 A1	21-11-1991
			ES 2063985 T3	16-01-1995
			JP 2606980 B2	07-05-1997
			JP 4227245 A	17-08-1992
WO 9825508	A	18-06-1998	US 5730758 A	24-03-1998
			AU 6936798 A	03-07-1998
			EP 1030587 A2	30-08-2000
			WO 9825508 A2	18-06-1998
US 5560532	A	01-10-1996	NONE	